

# Sandia National Laboratories / New Mexico

PROPOSAL FOR NO FURTHER ACTION
ENVIRONMENTAL RESTORATION PROJECT
SITE 46, OLD ACID WASTE LINE OUTFALL SITE
OPERABLE UNIT 1309

**June 1995** 

Environmental Restoration Project



United States Department of Energy Albuquerque Operations Office



GARY E. JOHNSON GOVERNOR

# State of New Mexico ENVIRONMENT DEPARTMENT

Underground Storage Tank Bureau
District I Office
4131 Montgomery, NE
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MARK E. WEIDLER SECRETARY

EDGAR T. THORNTON, III
DEPUTY SECRETARY

April 5, 1995

Jim Brinkman
Roy F. Weston
c/o Sandia National Laboratories
2501 Buena Vista S.E.
Mail Stop 1347
Department # 7582
Albuquerque, New Mexico 87185-1347

RE: NO FURTHER ACTION REQUIRED AT SANDIA NATIONAL LABORATORIES ENVIRONMENTAL RESTORATION SITE 40, OIL SPILL SITE, OPERABLE UNIT 1309, ALBUQUERQUE, NEW MEXICO

Dear Mr. Brinkman:

The New Mexico Environment Department reviewed the "Sandia Environmental Restoration Project, Site 40, Draft, No Further Action Proposal" submitted by Rarilee Conway on March 17, 1995. The Department determines that this site does not pose an immediate public health or environmental threat from the following reasons:

- data from soil borings drilled at the site show that soil contamination extends to 7 feet below ground surface,
- depth to ground water is 360 feet.

Therefore, the Department is not requiring additional work at this time. However, the Department reserves the right to require additional work in the future if data become available that indicate the presence of petroleum hydrocarbon contamination emanating from or in the vicinity of this site resulting in a threat to public health or the environment.

Thank you for your cooperation in this mater.

Sincerely,

Betsy Hovda Geologist

Underground Storage Tank Bureau

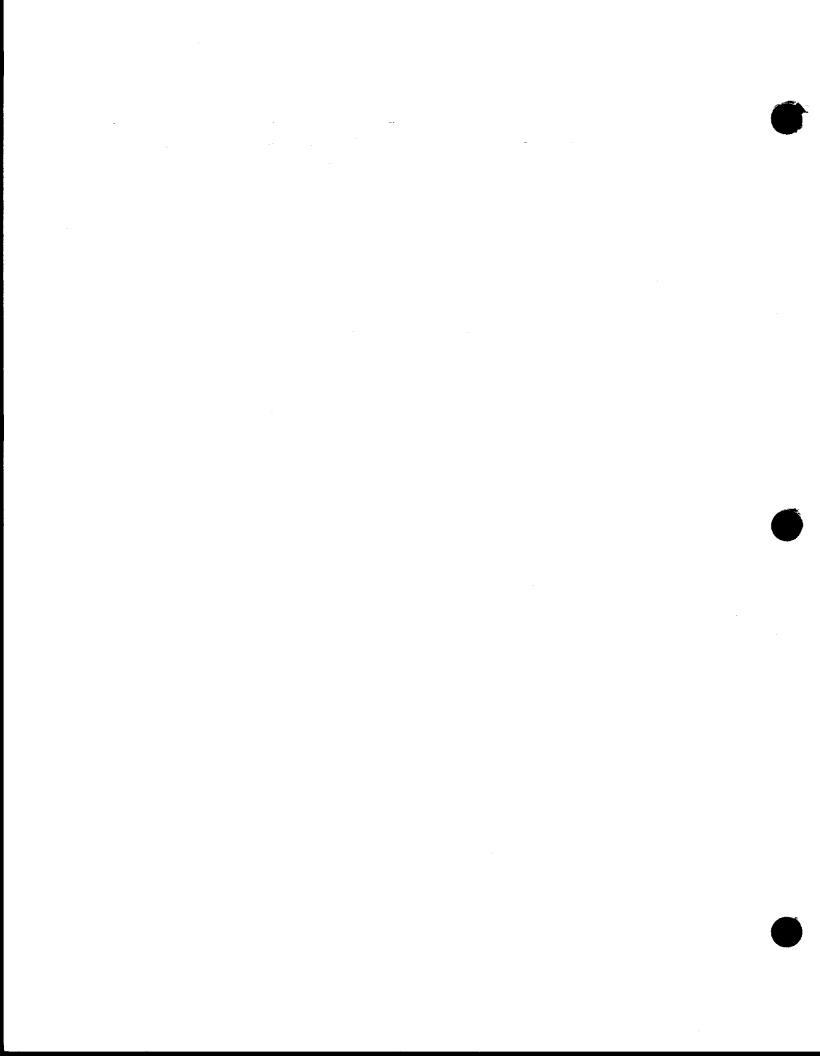
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# PROPOSAL FOR NO FURTHER ACTION

Site 46, Old Acid Waste Line Outfall Site Operable Unit 1309

SANDIA NATIONAL LABORATORIES/NEW MEXICO



# 1. Introduction

# 1.1 ER Site Identification Number and Name

Sandia National Laboratories/New Mexico (SNL/NM) is proposing a risk-based no further action (NFA) decision for Environmental Restoration (ER) Site 46, Old Acid Waste Line Outfall Site, Operable Unit (OU) 1309. ER Site 46 is listed in the Hazardous and Solid Waste Amendment (HSWA) Module IV (EPA August 1993) of the SNL/NM Resource Conservation and Recovery Act (RCRA) Hazardous Waste Management Facility Permit (NM5890110518) (EPA August 1992).

#### 1.2 SNL/NM Risk-Based NFA Process

This proposal for a determination of a risked-based NFA decision has been prepared using the criteria presented in Section 4.5.3 of the SNL/NM Program Implementation Plan (PIP) (SNL/NM February 1994). Specifically, this proposal will "contain information demonstrating that there are no releases of hazardous waste (including hazardous constituents) from solid waste management units (SWMU) at the facility that may pose a threat to human health or the environment" (as proposed in the code of Federal Regulations [CFR], Section 40 Part 264.51[a] [2]) (EPA July 1990). The HSWA Module IV contains the same requirements for an NFA demonstration:

Based on the results of the RFI [RCRA Facility Investigation] and other relevant information, the Permittee may submit an application to the Administrative Authority for a Class III permit modification under 40 CFR 270.42(c) to terminate the RFI/CMS [corrective measures study] process for a specific unit. This permit modification application must contain information demonstrating that there are no releases of hazardous waste including hazardous constituents from a particular SWMU at the facility that pose threats to human health and/or the environment, as well as additional information required in 40 CFR 270.42(c) (EPA August 1993).

For a risk-based proposal, an SWMU is eligible for an NFA determination if the NFA criterion established by the SNL/NM permit is met. This criterion, found in Section M.1 of the permit, is as follows: "[T]here are no releases of hazardous waste including hazardous constituents...that pose threats to human health and/or the environment..." This risk-base proposal contains information needed to make the NFA determination.

This proposal is using the technical approach which is the foundation for the SNL/NM corrective action process. The details of the SNL/NM technical approach are provided in Appendix C of the SNL/NM Program Implementation Plan (SNL/NM 1994). The first step in the technical approach is the data qualitative review step (the same step used to determine whether the SWMU is eligible for administrative NFA). Should significant uncertainties remain, the assessment of the SWMU continues with data collection.

At this site, sufficient data were not available to compare to established action levels or to develop site-specific action levels. Background soil samples were collected and analyzed to

develop upper tolerance limits (UTLs) for metals. Site-specific data were collected to compare to existing soil action levels (proposed Subpart S Action Levels) and UTLs. If site-specific concentrations exceeded the proposed Subpart S Action Levels or UTLs, then risk was analyzed. Site-specific soil concentrations were compared to the derived risk assessment action levels. Concentrations less than these action levels, either proposed Subpart S action levels, background UTLs, or derived risk-based values, triggered this NFA proposal for Site 46.

# 1.3 Local Setting

SNL/NM occupies 2,829 acres of land owned by the Department of Energy (DOE), with an additional 14,920 acres of land provided by land-use permits with Kirtland Air Force Base (KAFB), the United States Forest Service, the State of New Mexico, and the Isleta Indian Reservation. SNL/NM has been involved in nuclear weapons research, component development, assembly, testing, and other nuclear activities since 1945.

ER Site 46 (Figure 1) is located on land owned by DOE. The site is situated west and south of the Technical Area (TA) IV fence in a slight depression on top of the escarpment northwest of Tijeras Arroyo.

Surficial deposits in the SNL/KAFB area lie within four geomorphic provinces, which in turn contain nine geomorphic subprovinces. Site 46 lies within the Tijeras Arroyo subprovince. The Tijeras Arroyo subprovince is characterized by broad, west-sloping alluvial surfaces and the 50-meter-deep Tijeras Arroyo. The Tijeras Arroyo subprovince contains deposits derived from many sources, including granitic and sedimentary rocks of the Sandia Mountains, sedimentary and metamorphic rocks of the Manzanita Mountains, and sediments of the Upper Santa Fe Group.

# 2. History of the SWMU

# 2.1 Sources of Supporting Information

In support of this request for a risk-based NFA decision for ER Site 46, a background study was conducted to collect available and relevant site information. Interviews were conducted with SNL/NM staff and contractors familiar with site operational history.

The following information sources were available for the use in the evaluation of ER Site 46:

- Confirmatory-sampling program conducted in September 1994
- Risk analysis for three metals and three radionuclides
- One surface radiation survey
- One unexploded ordnance/high explosives (UXO/HE) survey
- Interviews and personnel correspondence
- Historical aerial photographs spanning 40 years
- Personal breathing zone air sampling for metals

# 2.2 Previous Audits, Inspections, and Findings

ER Site 46 was first listed as a potential release site based on the Comprehensive Environmental Assessment and Response Program (CEARP) interviews in 1985 (DOE September 1987). The Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) finding was uncertain for Federal Facility Site Discovery and Identification Findings (FFSDIF), Preliminary Assessment and Site Inspection; therefore, no Hazard Ranking System or Modified Hazard Ranking System migration mode scores were calculated for the SWMU (DOE September 1987). Site 46 was included in the Environmental Protection Agency (EPA) RCRA Facility Assessment (RFA) in 1987 (EPA April 1987).

# 2.3 Historical Operations

The Old Acid Waste Line carried wastes from several buildings in TA-I. It was installed between 1948 and 1950. The site begins as a north-south trending, 750-foot long open trench in a grassy field northwest of Building 981-1 in TA-IV. No pipe opening is visible at the "head" of the trench. As the trench crosses the field, it turns to the southeast and continues to an unengineered spillway above the Tijeras Arroyo floodplain. The spillway lies on a bank (40 to 50 feet of relief) composed of compacted alluvial sediment. Historical aerial photographs show vegetation, presumably supported by the discharge, growing southeast of the spillway to the active arroyo channel (about 200 feet distance from the spillway). The site is not restricted and is easily accessible.

During use, discharged effluent averaged an estimated 130,000 gallons per day. Use of the line was discontinued in the mid-to-late 1960s. The line received wastes from plating, etching, and photo processing operations; and cooling tower "blow down." Acids and metals are target contaminants. Chromic acid and ferric chloride are mentioned specifically in the site history, and ferric chloride was found in one soil sample collected during the RFA. Various radionuclides, possibly including tritium, uranium, and plutonium also were used in TA-I.

Building 863 (ER Site 98) was a source of discharge to the acid line. The information sheet for Site 98 indicates the presence of trichloromethane, silver, and photo processing chemicals with an ammonia-like odor. The waste solution from the silver recovery unit reportedly was discharged to the old acid waste line, which is the only specific information about chemical discharges.

# 3. Evaluation of Relevant Evidence

#### 3.1 Unit Characteristics

The Old Acid Waste Drainage Outfall is confined to the downstream natural drainage. All releases would be contained in this limited area. Most of the potential contamination resulting from discharged effluent would have most likely settled at or before the furthest extent of visible erosion/scour.

# 3.2 Operating Practices

The Old Acid Waste Drainage Outfall discharged industrial waste from TA-I from approximately 1948 until the mid-to-late 1960s. It has not been used since then.

# 3.3 Presence or Absence of Visual Evidence

The approximately 750-foot long trench is the only physical evidence of the outfall system. No discoloration of soils was observed during site reconnaissance and soil sampling activities.

# 3.4 Results of Previous Sampling/Surveys

In 1994, the site was visually surveyed for surface indications of UXO/HE. No UXO/HE were found (SNL/NM 1994a). Also, in 1994 a surface radiation survey was conducted on the entire site using an Eberline ESP-2 portable scaler, with an Eberline SPA-8 (2 inch X 2 inch sodium iodide) detector. A 30-second integrated count was performed at each proposed sample location, while scanning the detector over an area approximately 2 feet in radius around the sample location. The alarm was set at 1.3 times the background count rate. No alarms occurred during the survey. No surface anomalies were detected (SNL/NM 1994b).

# 3.5 Assessment of Gaps in Information

No environmental sampling data existed for Site 46. If contamination was present, potential constituents of concern (metals, radioactive constituents), would be expected at shallow depths. Metals and radioactive constituents generally adsorb on soil and precipitate rather than remaining soluble. A surface (0-6 inches deep) and shallow subsurface (6-36 inches deep) soil sampling program was developed and implemented in September 1994.

# 3.6 Confirmatory Sampling

The Confirmatory Sampling and Analysis Plan (SAP) can be found in Appendix A. Those soil sample results exceeding an action level are summarized in Table 1. A complete list of "hits" or detections and quality assurance (QA) results can be found in Appendix B.

For health and safety purposes, a photoionization detector, OVM, was used throughout the field program. The OVM measured no anomalous vapor concentrations.

Surface and shallow subsurface soil samples were collected at the most likely locations of contamination. Four samples were collected at the head of the site outfall (by the fire extinguisher training area west of TA-IV) and four samples were collected by the spillway into the Tijeras Arroyo drainage (Figure 1). Every sample was analyzed for metals<sup>1</sup>, chromium<sup>+6</sup>, total Kjeldahl nitrogen (TKN), and nitrate/nitrite. The four subsurface samples

Although the total analyte list (TAL) metal analytes include calcium, magnesium, potassium, and sodium, these nontoxic, major cations are not included in the evaluation. They do not pose a significant environmental or human health risk regardless of concentration.

also were analyzed for volatile organic compounds (VOCs). Four samples were analyzed for semivolatile organic compounds (SVOCs). As a general check for radioactive constituents, all the samples were analyzed for tritium, four samples were analyzed for isotopic uranium, and isotopic plutonium and all eight samples were screened in-house and two samples were screened off-site with gamma spectroscopy.

# 3.6.1 Background Samples for Metals and Radioactive Constituents

UTLs for background metals were calculated from analyses of 24 samples collected in the vicinity of the 11 sites discussed in the SAP (Appendix A). UTLs or background 95<sup>th</sup> percentiles for background radionuclides were calculated from samples collected throughout KAFB (IT 1994). A discussion of background calculations and supporting data and analyses are included in Appendices C and D.

# 3.6.2 Organic Compounds

No organic compounds were detected without qualification. Nitrate/nitrite was detected in seven of eight samples with concentrations ranging from 150 to 1400 milligrams per kilogram (mg/kg). TKN was detected in all eight samples, with concentrations ranging from 120 to 470 mg/kg. The main environmental or human health hazard pertaining to reduced nitrogen (as measured in TKN) is that it oxidizes to either nitrate or nitrite. Therefore, it is valid and conservative to compare TKN concentrations to action levels for nitrate and nitrite. The proposed Resource Conservation and Recovery Act (RCRA) Subpart S action levels for nitrate and nitrite are 100,000 mg/kg and 8,000 mg/kg, respectively. Cyanide was detected in two of eight samples, 46-01-B and 46-04-B, with concentrations of 0.16 and 0.18 mg/kg, respectively. The proposed Subpart S action level for cyanide is 2000 mg/kg. These results indicate no significant human health or environmental hazard because organic compounds only were detected tentatively and TKN, nitrate/nitrite and cyanide were detected at concentrations much lower than action levels.

#### 3.6.3 Metals

Personal breathing zone air sampling was used at Site 46 to monitor airborne particulate contamination for metals. No airborne metal contamination was detected. Selenium, mercury, and chromium<sup>16</sup> were not detected at Site 46. Silver was detected in two out of eight samples at concentrations of 0.59 and 0.58 mg/kg in Samples 46-01-B and 46-04-B, respectively. Silver was not detected in background samples. The proposed Subpart S action level is 400 mg/kg. Both tests comparing the site beryllium data to local background data indicated no contamination. The maximum local background value for beryllium was 0.53 mg/kg. Beryllium was not detected above 0.53 mg/kg at Site 46.

All other site metal concentrations, except for one analysis for cadmium, iron, and lead, were below the UTLs. For cadmium, Sample 46-01-B had a concentration of 4.0 mg/kg compared to a UTL of 3.82 mg/kg. The proposed Subpart S action level for cadmium in soils is 80 mg/kg. Sample 46-01-B had a concentration of 15 mg/kg for chromium, just exceeding the UTL of 14.3 mg/kg. However, the Subpart S action level for chromium is 80,000 mg/kg. For lead, Sample 46-04-B had a concentration of 27 mg/kg compared to a UTL of 23.1

mg/kg. A Subpart S action level was not proposed for lead. However, a memorandum from an EPA assistant administrator to EPA regional division directors does supply a risk-based action level for lead in soils, 400 ppm (mg/kg) (EPA 1994). This risk-based action level presumes that lead be considered individually rather than in conjunction with other metals. This action level of 400 mg/kg far exceeds the concentration in Sample 46-04-B of 27 mg/kg. For iron, Sample 46-02-A had a concentration of 17,000 mg/kg compared to a UTL of 16,962 mg/kg.

## 3.6.4 Radionuclides

Potassium-40 was detected in two samples at activities of 16.4 and 22.3 picocuries per gram (pCi/g), compared with the base-wide background UTL of 25.34 pCi/g. Lead-212 was detected in Sample 46-03-A at an activity of 0.89 pCi/g, compared to a base-wide background UTL of 1.0795 pCi/g (IT 1994). Plutonium-239/240 and plutonium-238 were not detected above the minimum detectable activity (MDA) in Site 46 samples. Uranium-238, uranium-235/236, and uranium-234 were detected in four samples at low activities below the base-wide background UTL and the maximum activity of six local background analyses. Thallium was not detected at Site 46. These constituents require no further evaluation.

Lead-214 was detected in Sample 46-03-A at an activity of 0.93 pCi/g, compared with the base-wide background UTL of 0.90 pCi/g. Radium-226 was detected in Samples 46-02-B, 46-03-B and 46-04-B with activities of 2.74, 2.14, and 2.06 pCi/g, respectively. The base-wide background UTL for radium-226 is 1.94 pCi/g (IT 1994). Additional off-site radiological analyses for radium-226 were requested for Samples 46-01-A, 46-01-B, 46-02-B, 46-03-B, and 46-04-B. These results indicated activities less than 2.74 pCi/g. Tritium was detected in all eight samples at activities ranging from 0.023 to 0.17 pCi/g; tritium was not detected above the MDA in local background samples.

# 3.6.5 Quality Assurance Results

As discussed in the Confirmatory Sampling and Analysis Plan (Appendix A), quality assurance samples, including field duplicates, trip blanks and rinsates, were collected as part of the 11 site sampling program. Analyses indicate that the field soil duplicates were comparable to the original soil sample results. The trip blanks and rinsates indicated no significant sampling contamination. QA results can be found in Appendix B. Level I and Level II data verification was conducted on all data, as described in the PIP (SNL/NM 1994).

# 3.7 Risk Analysis

To further evaluate the data for metals with concentrations greater than background UTLs, a risk assessment was performed for a combination of cadmium, iron, and silver, assuming the maximum detected concentrations. To further evaluate the site data for radionuclides with activities above background UTLs or those without background UTLs, a risk analysis was performed for the combination of lead-214, tritium, and radium-226, assuming the maximum detected activities.

The risk calculations were designed to produce conservatively large estimates of hazard index and radioactive dose to counter uncertainties in the soil data. This approach facilitates the following decision regarding future activities at Site 46:

- If the conservative estimates based on the soil data result in an unacceptable hazard index (greater than 1) or dose (greater than 10 mrem/year), further investigation and/or remediation will be needed; or
- If the hazard index and dose estimates are acceptable, the potential for health hazards at the site is extremely low, and further actions will not be needed.

Hazard indices and radionuclide doses were computed using methods and equations promulgated in proposed RCRA Subpart S documentation (EPA 1990). Accordingly, all calculations were based on the assumption that receptor doses from both toxic metals and radionuclides result from ingestion of contaminated soil.

Calculation of hazard indices required values of oral reference doses (oral RfDs) for each of the metals. The RfD values for cadmium and silver were taken from EPA's IRIS database (IRIS 1994). The RfD for iron is a provisional value provided by EPA Region VI personnel.

Similarly, calculation of radionuclide doses required values of dose conversion factors, which are used to convert radionuclide intakes (in units of pCi/year) into effective dose equivalents (in units of mrem/year). Published values of dose conversion factors (Eckerman et al., 1988 and Gilbert et al., 1989) exist for lead-214, tritium, and radium-226.

To assure that the computed hazard indices and doses were conservatively large, only the maximum observed concentration of each constituent at a site was employed. To consider combined effects, a hazard index was calculated as the sum of the individual metal hazard quotients and a radiological dose was calculated as the sum of the individual doses.

Following proposed Subpart S methodology, the equation and parameter values used to calculate the summed hazard index for toxic metals were:

$$HI = \sum_{i} [HSR(i) \times S(i)]$$

**(1)** 

where:

HI = total hazard index (dimensionless),

HSR(I) = hazard index-to-soil concentration ratio for the i<sup>th</sup> metal (kg/mg)

$$= \frac{1 \times A}{RfD(i) \times W} \times \frac{0.001 \text{ g}}{mg}$$

S(I) = soil concentration of the i<sup>th</sup> metal (mg/kg), I = soil ingestion rate = 0.2 g/day, A = absorption factor (dimensionless) = 1, W = body weight = 16 kg, and RfD(I) = oral reference dose for the i<sup>th</sup> metal (mg/kg-day).

Risk assessment guidance, prepared by the EPA (EPA 1989), recommends that the total hazard index be less than one in order for a site to be considered a non-threat to human health.

Following proposed Subpart S methodology, the equation and parameter values used to calculate the summed radioactive dose were:

DOSE = 
$$\sum_{i}$$
 [DSR(i) x S(i)]

(2)

where:

DOSE	=	total effective dose equivalent (mrem/yr);
DSR(I)	=	dose-to-soil concentration ratio for the ith radionuclide
		(mrem/yr)/(pCi/g), = I X DCF(I);
S(I)	=	soil concentration of the i <sup>th</sup> radionuclide (pCi/g);
I	==	soil ingestion rate = $0.2 \text{ g/day} = 73 \text{ g/yr}$ ; and
DCF(I)	=	dose conversion factor for the i <sup>th</sup> radionuclide (mrem/pCi).

The PIP stipulates that, for the purpose of computing media action levels, the total radioactive dose at a site should not be greater than 10 mrem/year (DOE 1994), which corresponds to a cancer risk of less that 10<sup>-6</sup> excess deaths.

The input and results of the risk calculations are presented in Tables 2 and 3. The summed hazard index for metals is less than 1 and the summed radioactive dose is less than 10 mrem/year. Therefore, the site is considered to be risk-free in terms of metals and radionuclide contamination.

# 3.8 Rationale for Pursuing a Risk-Based NFA Decision

In September, 1994, surface soil and shallow subsurface soil samples were collected at the "head" of the trench (where the flow spills into the natural drainage) and at the furthest extent of visible erosion/scour where the discharged effluent would have most likely settled. These

two areas are the most likely areas for contamination. SNL/NM is proposing a risk-based NFA because representative soil samples from ER Site 46 have concentrations less than action levels; either proposed Subpart S action levels, background UTLs, background 95<sup>th</sup> percentiles, or derived risk-based values.

# In addition

- A site visit in 1993 by ER personnel confirmed the presence of a confined natural drainage with no discoloration in the soils.
- In June 1994, a UXO/HE visual survey was conducted by KAFB Explosive Ordnance Division (EOD) and found no UXO/HE ordnance debris at Site 46 (SNL/NM 1994a).
- In September, 1994, as part of the surface soil sampling effort at Site 46, a surface radiation survey was conducted (SNL/NM 1994b). No surface anomalies were detected at Site 46.
- In September 1994, as part of the surface soil sampling effort at Site 46, particulate metal contamination was monitored with Personal Breathing Zone Air Sampling. No airborne contamination was detected.

# 4. Conclusion

Based upon the evidence cited above, ER Site 46 has no releases of hazardous waste or hazardous constituents that pose a threat to human health and/or the environment. Therefore, ER Site 46 is recommended for an NFA determination.

#### 5. References

# 5.1 ER Site References

Eckerman, K.F., A.B. Wolbarst, and A.C.B. Richardson, 1988. "Limiting Values of Radionuclide Intake and Air Concentration and Dose Conversion Factors for Inhalation, Submersion, and Ingestion," EPA -520/1-88-020, Federal Guidance Report No. 11, prepared by Oak Ridge National Laboratory, Oak Ridge, TN, for Office of Radiation Programs, Washington, DC.

Gilbert, T.L., C. Yu, Y.C. Yuan, A.J. Zielen, M.J. Jusko, and A. Wallo, 1989. Implementing Residual Radioactive Material Guidelines, A Supplement to U.S. Department of Energy Guidelines for Residual Radioactive Material at Formerly Utilized Sites Remedial Action Program and Surplus Facilities Management Program Sites, prepared by Argonne National Laboratory for U.S. Department of Energy, ANL/ES-160, DOE/CH/8901, 203 pp.

Guttman, I., 1970. Statistical Tolerance Regions: Classical and Bayesian, Hafner Publishing, Darien Connecticut.

IRIS, Integrated Risk Information System (data base), 1994. U.S. Environmental Protection Agency, Office of Research and Development.

International Technology (IT) Corporation, 1994. Draft "Background Concentrations of Constituents of Concern to the Sandia National Laboratories/New Mexico Environmental Restoration Project Phase II: Interim Report."

Sandia National Laboratories/New Mexico (SNL/NM), 1994a. "Unexploded Ordnance/High Explosives (UXO/HE) Visual Survey of ER Sites Final Report," Sandia National Laboratories, Albuquerque, New Mexico."

Sandia National Laboratories/New Mexico (SNL/NM), 1994b. "Summary of Radiological Survey Results by SNL Dept. 7714 for ER Sites 50, 227, 229, 230-234," Sandia National Laboratories, Albuquerque, New Mexico.

U.S. Environmental Protection Agency (EPA), July 1994. Guidance on Residential Lead-Base Paint, Lead-Contaminated Dust, and Lead-Contaminated Soil, memorandum from Lynn R. Goldman, M.D., EPA Assistant Administrator to EPA Division Regional Directors.

U.S. Environmental Protection Agency (EPA), July 1990. "Corrective Action for Solid Waste Management Units (SWMU) at Hazardous Waste Management Facilities, Proposed Rule," *Federal Register*, Vol. 55, Title 40, Parts 264, 265, 270, and 271.

U.S. Environmental Protection Agency (EPA), 1989. "Statistical Analysis of Ground-Water Monitoring Data at RCRA Facilities, Addendum to Interim Final Guidance," U.S. Environmental Protection Agency, Office of Solid Waste, Waste Management Division, Washington DC.

# 5.2 Reference Documents

Department of Energy (DOE), September 1987. "Comprehensive Environmental Assessment and Response Program, Phase I Installation Assessment Sandia National Laboratories - Albuquerque," Department of Energy Albuquerque Operations Office, Environmental Safety and Health Division, Environmental Program Branch, September 1987.

Sandia National Laboratories/New Mexico (SNL/NM), August 1994. Environmental Restoration Project Information Sheet for Site 46, Storm Drain System Outfall, Sandia National Laboratories, Albuquerque, New Mexico.

U.S. Environmental Protection Agency (EPA), April 1987. "Final RCRA Facility Assessment Report of Solid Waste Management Units at Sandia National Laboratories, Albuquerque, New Mexico," Contract No. 68-01-7038, EPA Region VI.

Sandia National Laboratories/New Mexico (SNL/NM), February 1994. Draft "Program Implementation Plan for Albuquerque Potential Release Sites," Sandia National Laboratories, Albuquerque, New Mexico.

- U.S. Environmental Protection Agency (EPA), August 1993, "Module IV of RCRA Permit No. NM 5890110518, EPA Region VI, issued to Sandia National Laboratories, Albuquerque, New Mexico.
- U.S. Environmental Protection Agency (EPA), August 1992, "Hazardous Waste Management Facility Permit No. NM5890110518, EPA Region VI, issued to Sandia National Laboratories, Albuquerque, New Mexico.
- U.S. Environmental Protection Agency (EPA), April 1987, "Final RCRA Facility Assessment Report of Solid Waste Management Units at Sandia National Laboratories, Albuquerque, New Mexico," Contract No. 68-01-7038, EPA Region VI.

# 5.3 Aerial Photographs

Ebert & Associates, Inc., November 1994. "Photo-Interpretation and Digital Mapping of ER Sites 7,16,45,228 from Sequential Historical Aerial Photographs."

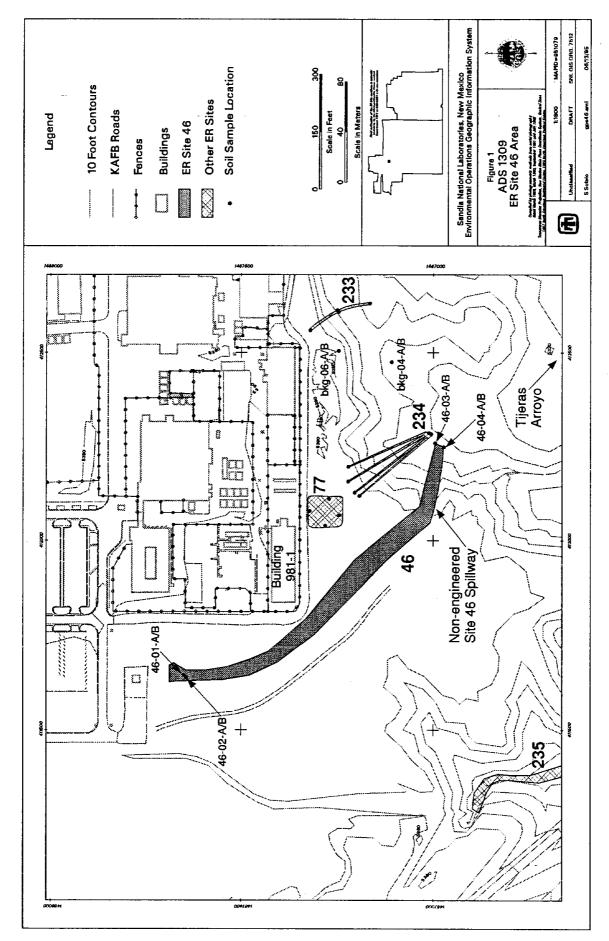


Figure 1. Old Acid Waste Line Outfall Site 46.

Table 1. Site 46 - Results of Shallow Soil Sampling and Analysis

Sample Identifier	Analytical Method	Constituent	Concentration (mg/kg)	Qualifier(s)	Background (mg/kg)	Action Level(s) (mg/kg)
46-01-B	VOCs (8240)	2-butanone	0.003	JВ		
46-02-B	VOCs (8240)	2-butanone	0.005	JB	***	
46-03-B	VOCs (8240)	2-butanone	0.005	JB		
46-04-B	VOCs (8240)	2-butanone	0.004	JB		
46-01 <b>-</b> B	SVOCs (8270)	Di-n-butyl phthalate	0.066	J		
46-01-B	TAL Metals (6010)	Cadmium	4.		3.82	80/5.1*
46-02-A	TAL Metals (6010)	Iron	17000.		16,962	21500*
46-01-B	TAL Metals (6010)	Silver	0.59			400/7.5*
46-01-B	TAL Metals (6010)	Silver	0.58			400/7.5*
46-01-B	TAL Metals (6010)	Chromium	15		14.3	80,000
46-04-B	TAL Metals (6010)	Lead	27		23.1	400**
46-01-B	Cyanide (acid digestion)	Cyanide	0.16			2,000
46-04-B	Cyanide (acid digestion)	Cyanide	0.18	<u> </u>		2,000
46-01-A	TKN (acid digestion)	TKN	310.			100,000/8,000
46-01-B	TKN (acid digestion)	TKN	380.			100,000/8,000
46-02-A	TKN (acid digestion)	TKN	360.			100,000/8,000
46-02-B	TKN (acid digestion)	TKN	470.	<del></del>		100,000/8,000
46-03-A	TKN (acid digestion)	TKN	120.			100,000/8,000
46-03-B	TKN (acid digestion)	TKN	130.			100,000/8,000
46-04-A	TKN (acid digestion)	TKN	160.			100,000/8,000
46-04-B	TKN (acid digestion)	TKN	190.			100,000/8,000
46-01-A	NO3/NO2 (353.2)	NO3/NO2	1000.			100,000/8,000
46-01-B	NO3/NO2 (353.2)	NO3/NO2	1230.			100,000/8,000
46-02-A	NO3/NO2 (353.2)	NO3/NO2	1300.			100,000/8,000
46-02-B	NO3/NO2 (353.2)	NO3/NO2	1200.			100,000/8,000
46-03-A	NO3/NO2 (353.2)	NO3/NO2	1400.			100,000/8,000
46-03-B	NO3/NO2 (353.2)	NO3/NO2	150.			100,000/8,000
46-04-A	NO3/NO2 (353.2)	NO3/NO2	410.	****		100,000/8,000
46-03-A	Gamma Spec (Off-site)	Lead-214	0.93 pCi/g		0.9 pCi/g	42.2 pCi/g*
46-01-A	Tritium (600 906.0)	Tritium	0.038 pCi/g		,	7.7 pCi/g*
46-01-B	Tritium (600 906.0)	Tritium	0.038 pCi/g			7.7 pCi/g*
46-02-A	Tritium (600 906.0)	Tritium	0.044 pCi/g	<del></del>		7.7 pCi/g*
46-02-B	Tritium (600 906.0)	Tritium	0.039 pCi/g			7.7 pCi/g*
46-03-A	Tritium (600 906.0)	Tritium	0.023 pCi/g		***	7.7 pCi/g*
46-03-B	Tritium (600 906.0)	Tritium	0.04 pCi/g			7.7 pCi/g*
46-04-A	Tritium (600 906.0)	Tritium	0.026 pCi/g			7.7 pCi/g*
46-04-B	Tritium (600 906.0)	Tritium	0.17 pCi/g			7.7 pCi/g*
46-02-B	Gamma Spec (In-house)	Radium-226	2.74 pCi/g	1	1.94 pCi/g	124 pCi/g*
46-03-B	Gamma Spec (In-house)	Radium-226	2.14 pCi/g		1.94 pCi/g	124 pCi/g*
46-04-B	Gamma Spec (In-house)	Radium-226	2.06 pCi/g		1.94 pCi/g	124 pCi/g*

# Table 1. Site 46- Results of Shallow soil Sampling and Analysis (Concluded)

# <u>Notes</u>

"J" qualifier means detected at a concentration below the laboratory reporting limit.

"B" qualifier means detected in the associated blank sample.

For the metals, background is the 95 percent upper tolerance level for the local background data.

For lead-214 and radium-226, background is the 95 percent upper tolerance level for the base-wide background data.

Action levels without an asterisk are proposed Subpart S Action Levels.

Proposed Subpart S action levels for nitrate and nitrite are 100,000 and 8,000 mg/kg, respectively.

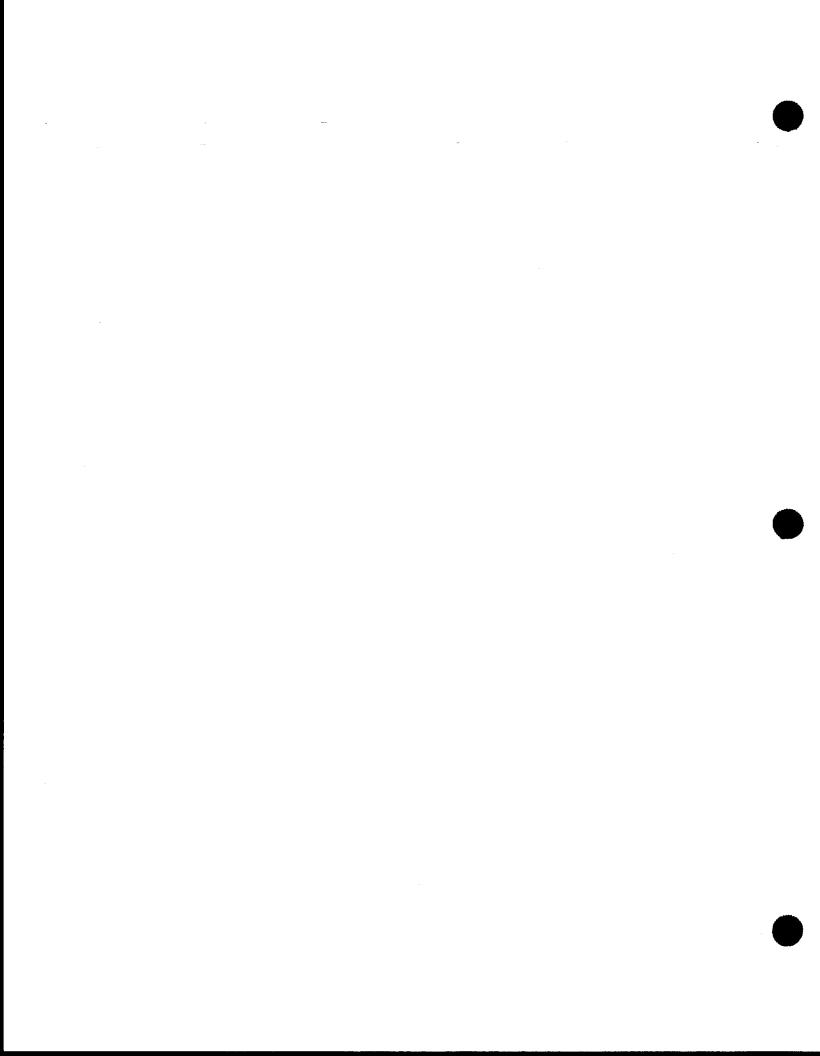
Action levels followed by one asterisk are calculated risk-based levels.

Table 2. Metal Risk Calculations for Site 46

Constituent	Concentration (mg/kg)	RfD(I) (mg/kg-day)	Individual HI	Source of RfD						
Iron	17000.	3.00E-01	7.08E-01	Provisional RfD provided by EPA Region VI.						
Cadmium	4.00E+00	1.00E-03	5.00E-02	IRIS						
Silver	5.90E-01	5.00E-03	1.48E-03	IRIS						
Summed HI		·	7.60E-01	77 P						

Table 3. Radionuclide Risk Calculations for Site 46

Constituent	Activity (pCi/g)	DCF(I) (mrem/pCi)	Individual Dose (mrem/year)	Source of DCF Eckerman et al., 1988				
Lead-214	9.30E-01	7.80E-06	5.30E-04					
Radium-226	2.74E+00	1.10E-03	2.20E-01	Gilbert et al., 1989				
Tritium	1.70E-01	6.30E-08	7.82E-07	Gilbert et al., 1989				
Summed Dose		<u> </u>	2.21E-01	144				



# APPENDIX A

Confirmatory Sampling and Analysis Plan

# APPENDIX B

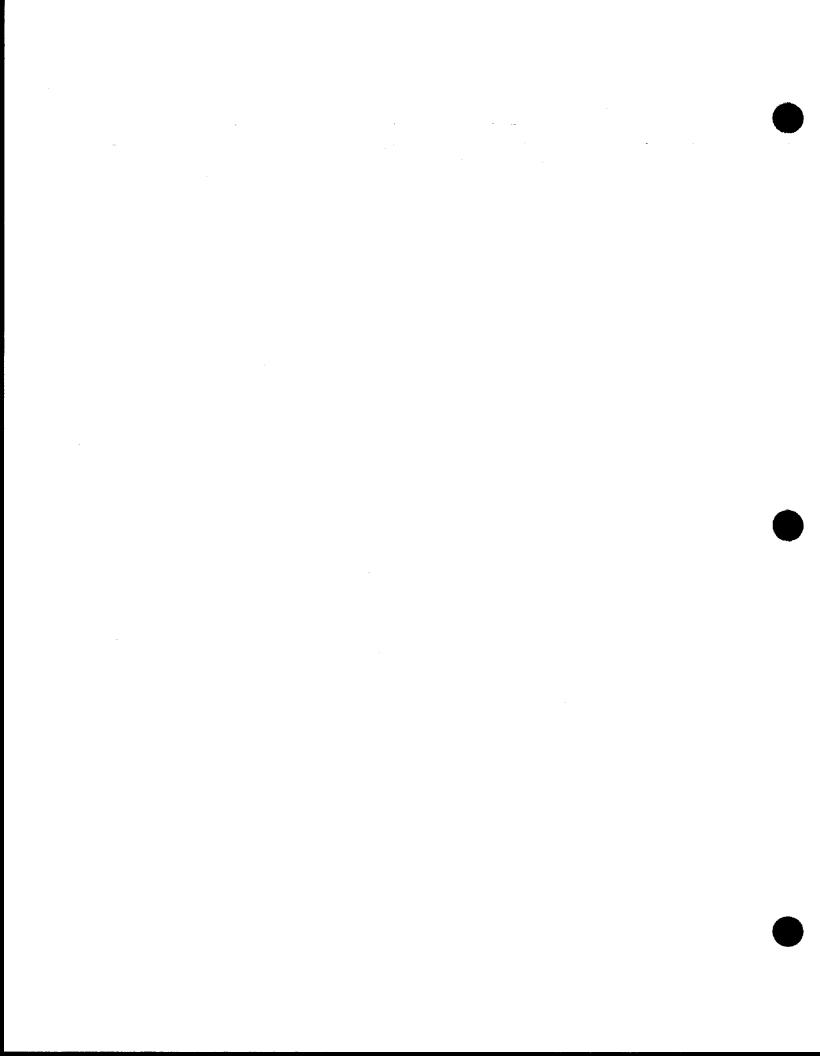
**Analytical Results** 

# APPENDIX C

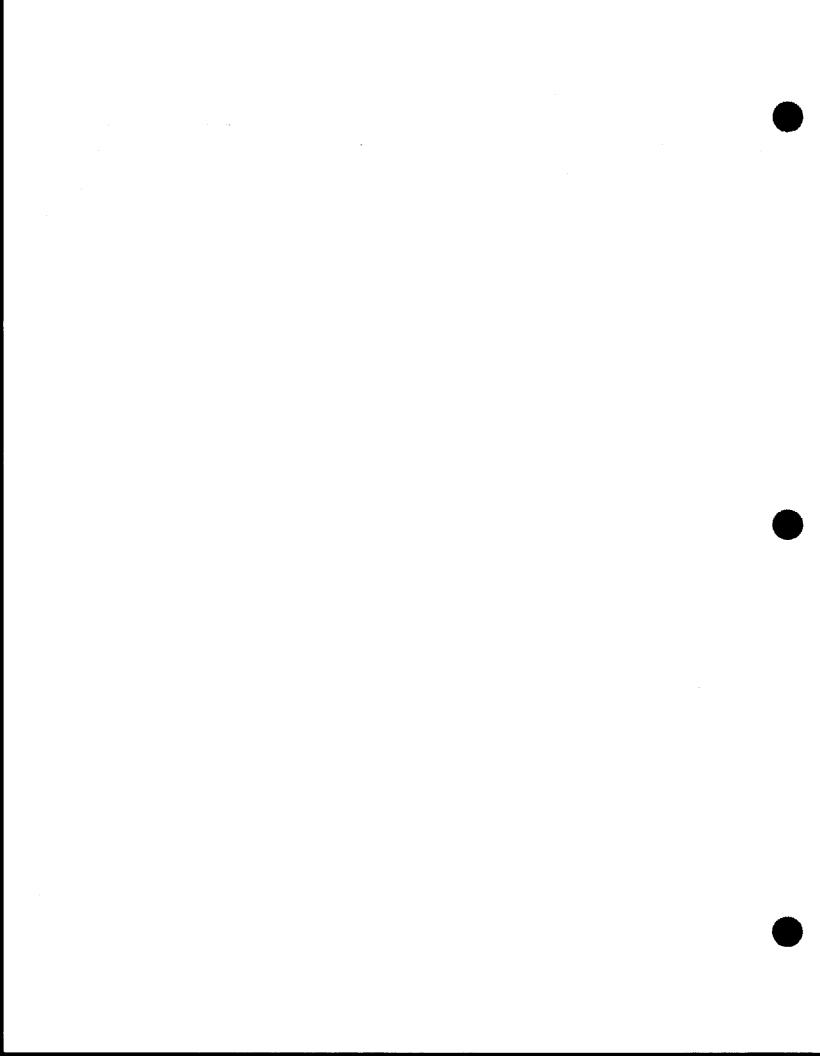
Background Calculations for Metals and Radionuclides

# APPENDIX D

Probability Plots, Local Background UTL Calculations, and Base-wide Background UTLs for Radionuclides



# Appendix A Confirmatory Sampling and Analysis Plan



# SAMPLING AND ANALYSIS PLAN FOR ELEVEN SITES IN TIJERAS ARROYO OPERABLE UNIT SANDIA NATIONAL LABORATORIES/ NEW MEXICO

#### Introduction

The purpose of the sampling and analysis described in this plan is to determine the appropriate way to proceed toward closure of 11 ( of the 17) sites in the Tijeras Arroyo Operable Unit. Based on the surface and shallow subsurface soil samples and analyses for the constituents of concern (COCs), one of three approaches will be pursued for each site:

- A petition for "No Further Action" (NFA) will be produced for regulatory consideration;
- 2. A voluntary corrective measure (VCM) will be designed and implemented, hopefully followed by an NFA petition; or
- 3. The site assessment and eventual closure will follow the standard RFI/CMS path

Most of the sites covered by this Sampling and Analysis Plan (SAP) are outfalls from the storm water and sanitary sewer systems emanating from Sandia Technical Areas (TAs) I, II, and IV. The general sampling program for the outfalls will be to collect four samples at the head of the outfall, two samples of surface soil (0 to 6 inches deep) and two samples of shallow subsurface soil (18 to 36 inches deep) and four samples (two surface soil and two shallow subsurface soil) at the furthest extent of channel erosion and scour. The analytes for most of the samples are volatile organic compounds, semi-volatile organic compounds (BNAs), metals, chromium<sup>+6</sup>, for samples where chromium is found in a metals analysis, total petroleum hydrocarbon (TPH), explosives, Total Kjeldahl Nitrogen (TKN), nitrate/nitrite, and Gamma Spectroscopy for radionuclides, isotopic uranium, isotopic plutonium, tritium, and chlorodiphenyls (PCBs).

#### Sampling Procedures and Volumes

Surface soil samples will be collected with a stainless steel scoopula or trowel and placed in a stainless steel bowl. After at least 1000 ml<sup>1</sup> of soil has been collected, the soil will be thoroughly mixed in the bowl and transferred to two or three 500-ml sample bottles with a stainless steel scoopula. Sample bottles will be labeled accordingly and the appropriate sample information (sample depth, collection date and time, etc.) will be documented on the chain-of custody (COC) after each sample is collected. Samples will then be packaged and cooled to 4 degrees Celsius.

Shallow subsurface soil samples (18-36 inches) will be collected with a 2-inch (minimum) hand auger. A soil sample is collected by turning the auger clockwise and advancing it into the ground until the bucket at the end of the auger (last 6-8 inches) is full of soil or refusal occurs. Several runs with the auger is anticipated in order to obtain the appropriate volume. A hand shovel may also be used to bypass large rocks in order to continue with the auger. The auger is then extruded counter-clockwise from the ground and the soil is removed from the auger and placed in a stainless steel bowl. After 1,125² ml of soil has been collected, the soil will be mixed in the bowl and transferred to two or three 500-ml sample bottles and one 125-ml sample bottle with a stainless steel scoopula. Sample bottles will be labeled accordingly and the appropriate sample information will be documented on the COC after each sample is collected. Samples will then be packaged and cooled to 4 degrees Celsius.

#### Waste Generation and Equipment Decontamination

Decontamination of sampling equipment will be done between each sample.

Decontamination will include thoroughly washing the inside and outside of the sampling equipment with a spray of ALCONOX™ or LIQUINOX™ and water; rinsing with distilled,

<sup>&</sup>lt;sup>1</sup>The sample volume varies between 1,000 and 1,500 ml depending on the analyses for the sample.

<sup>&</sup>lt;sup>2</sup>The sample volume varies between 1,125 and 1,625 ml depending on the analyses for the sample.

deionized water; and drying before reusing. No soil waste will be generated. The soil removed from the hand-auger holes, while collecting samples at a depth of 18 to 36 inches, will be return to the hole. The sampling tools, which are scoopulas/trowels, hand-augers, and shovels, will be decontaminated with water and ALCONOX™ after each use. The decon leachate will be stored in capped 1-gallon containers. One or two containers will be used for each site and two to four containers will be used for the background samples. The containers will be labeled as "IDW" and the site number identified on each container. All the containers will be stored at Site 232, a central location. The leachate waste will be disposed according to the analytical results of the soil samples collected at the site.

#### Site Descriptions

The sites that will be sampled are

- Site 46, Old Acid Waste Line Outfall;
- Site 50, Old Centrifuge Site;
- Site 77, Oil Surface Impoundment;
- Site 227, Bldg. 904 outfall;
- Site 229, Storm Drain System Outfall;
- Site 230, Storm Drain System Outfall;
- Site 231, Storm Drain System Outfall;
- Site 232, Storm Drain System Outfall:
- Site 233, Storm Drain System Outfall;
- Site 234, Storm Drain System Outfall; and
- Site 235, Storm Drain System Outfall.

The site locations are shown in Figure 1. A description of the site history, conditions, previous investigations, and sampling plans are described in the following sections.

#### Site 46: Acid Waste Line Outfall

The Old Acid Waste Line carried wastes from several buildings in TA I. The waste line begins as a north-south trending, 750-feet long open trench in a grassy field northwest of Building 981-1 in TA IV. No pipe opening is visible at the "head" of the trench. As the trench crosses the field, it turns to the southeast and continues to a non-engineered spillway at the edge of Tijeras Arroyo. The spillway lies on a bank (40 to 50 feet of relief) composed of compacted alluvial sediment. Historical aerial photographs show vegetation, presumably supported by the discharge, growing southeast of the spillway to the active arroyo channel (about 200 feet distance from the spillway). The site is not restricted and is easily accessible.

During use, discharged effluent averaged an estimated 130,000 gallons per day. Use of the line has been discontinued. The line received wastes from plating, etching, and photo processing operations, and cooling tower "blow down". Acids and metals are target contaminants. Chromic acid and ferric chloride are mentioned specifically in the site history, and ferric chloride was found in the soils during a limited sampling event. Various radionuclides, possibly including tritium, uranium, and plutonium were used in TA I.

Building 863 was a source of discharge to the Acid Line. The information sheet for ER Site 98 (Building 863, TCA Photochemical Release: Silver Catch Boxes) indicates the presence of trichloromethane, silver, and photo-processing chemicals with an ammonia-like odor. The waste solution from the silver recovery unit reportedly was discharged to the Old Acid Waste Line, which is the only specific information about chemical discharges.

The site has been visually surveyed for surface indications of unexploded ordnance and high explosives (UXO/HE). No UXO/HE were found. Also, a surface radiation survey was

conducted on the entire site. No surface radiation anomalies were detected.

The sampling program includes four samples collected at the "head" of the site outfall (by the fire extinguisher training area west of TA IV) and four samples collected by the spillway into the Tijeras Arroyo drainage (Figure 1). Every sample will be analyzed for tritium, metals, chromium<sup>+6</sup> (if chromium is detected), TKN, and nitrate/nitrite. Half the samples will also be analyzed for semi-volatiles and cyanide. Additionally, all the subsurface samples will be analyzed for volatiles. The analytes are listed in Table 1. A "4" on the table indicates that ALL the samples will be analyzed

for that specific analyte whereas a "2" on the table indicates half the samples will have additional analyses for the analyte listed.

#### Site 50: Old Centrifuge

Site 50, Old Centrifuge, was an outdoor, rocket propelled centrifuge that was used in the early 1950s to test units under G forces. The facility is located east of the TA II fence in a slight depression on top the escarpment northwest of Tijeras Arroyo. The concrete centrifuge pad has a diameter of 80 to 90 feet. The site has a 7-foot high wooden retaining wall on the north, east, and south sides. The west side is open. The centrifuge arm assembly, which has a 20-foot radius, is sitting outside the wall to the north and appears to be intact. Control wiring to the center axis of the centrifuge was suspended from a cable between two telephone poles on the north and south side of the pad. The control wiring went to a bunker located to the southwest over the escarpment. The bunker had a electrical transformer containing PCB. The electrical transformer has been removed. The pad was not stained and no spills or leaks were reported.

The centrifuge was rocket driven by two T40 6-KS-3000 or two Deacon 3.5DS-5700 solid rocket motors. The combustion byproducts produced by these rocket motors were carbon dioxide, carbon monoxide, water, hydrochloric acid, aluminum oxide, and possibly barium oxide. No other HE is known or suspected at the site. The rocket orientation would expel combustion byproducts towards the retaining wall and the opening to the west. The rocket propellant would be consumed in the rocket motor case. Under normal operating conditions, no unburned propellant would be released.

In 1987, a reconnaissance investigation at five potential contaminated sites, including the Old Centrifuge Site, was conducted by the ER Project. Samples were analyzed for uranium, TNT, HSL inorganics, TCLP constituents, and EP Toxicity constituents. Metals, including barium, were detected at concentrations well below regulatory action levels. Total uranium concentrations were typical of area background levels. TNT, pesticides, PCBs, herbicides, and semi-volatiles TCLP compounds were not detected.

Prior to sampling, the surface will be surveyed for radiation. If contamination exists, it is expected to be around the edge of the centrifuge pad at the surface, probably along the open west side. The constituents of concern are metals (specifically lead, beryllium, and barium), depleted uranium, and high explosives. Four surface samples and four subsurface samples will be collected. The sampling locations will be biased toward the west side of the site because that is the open side (Figure 1). All surface samples will be analyzed for all the COCs. One-half of the subsurface samples will be analyzed for uranium and high explosives. All four subsurface samples will be analyzed for metals.

#### Site 77: Oil Surface Impoundment

The Oil Surface Impoundment Site is outside the TA IV fence, southeast of Building 981-1. The surface impoundment, which was constructed in the 1970's, is used to catch waste water from accelerators. At the time of the RCRA facilities environmental survey, the impoundment was unlined. Since then the impoundment was drained. Soil samples were analyzed for PCBs and

solvents. Based on the analytical results, the impoundment was determined to be clean. Subsequently, the impoundment was lined with geotextile and is now regulated under Sandia's Surface Water Discharge Program.

This site will not require UXO/HE or radiation surface surveys. Minimal confirmation sampling and analysis is proposed to verify that the site is clean. Three surface and three shallow subsurface samples are proposed. The samples will be collected along the perimeter of the existing lined pond (Figure 1). All the samples will be analyzed for PCBs. The subsurface soil samples also will be analyzed for volatile organic compounds (Table 1).

# Site 227: Bunker 904 Outfall

Site 227 is an inactive outfall from the septic system for Building 904 (ER Site 48) in TA II. The site starts where the discharge exits the septic tank piping system, approximately 100 feet northeast of the southernmost point of TA II. The extent of the area influenced by the discharge may include the bank of Tijeras Arroyo below the outfall and some area between the outfall and the main channel of Tijeras Arroyo. The site is along the eastern edge of ER Site 45.

Building 904, built in 1948, was used for weapons assembly, HE testing, photo processing, and various other testing. Sanitary wastes were discharged to a septic tank, and other wastes were discharged to the outfall.

Mineral oil is also being considered a potential soil contaminant at all outfalls along the Tijeras Arroyo due to a recent release (June 1994) of mineral oil at Outfall 232 and vague historical records.

Possible soil contaminants are explosives, radioactive materials from weapons processing, including tritium, uranium, and plutonium, solvents (acetone, methylene chloride, methyl ethyl ketone, carbon tetrachloride, toluene, xylene, hexane, alcohols), and inorganics (ammonium hydroxide, barium, cadmium, silver, chromium, titanium, cyanide).

Access to this site is along the TA II perimeter road. This site is within the TA II testing exclusion zone. The best days to sample are generally Friday, Saturday, and Sunday, when testing ceases. Bruce Berry (telephone 845-8018) must be contacted to gain permission and access to this site. Prior to sampling

- 1. tumbleweeds will be cleared from locations to be sampled and placed adjacent to the drainage;
- 2. these locations will be visually scanned for UXO/HE, and
- 3. these locations will be screened for surface radiation anomalies.

The proposed sampling program is to collect four surface soil samples and four shallow subsurface samples. Two surface and two subsurface samples will be collected at the outfall. The other two surface and two subsurface samples will be collected at the furthest visible channel erosion and scour (Figure 1). The analytes are listed in Table 1.

# Sites 229 - 235: Storm Drain Systems Outfalls

These sites consist of the discharge areas at seven outfalls along the northern embankment of Tijeras Arroyo. The outfalls discharged industrial effluent and storm water from TAs I, II, and IV. Presently they only discharge storm water. The outfalls receive runoff from Site 96 (Storm Drain System) and other engineered drain systems within the three TAs. The sites are along approximately 3/2 miles of the embankment.

The specific constituents in the industrial effluent at these sites are not known. The possible discharged contaminants include chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, and other petroleum products. To cover this array of possible contaminants, soil samples will be analyzed for volatiles (subsurface samples only), semi-volatiles, metals and chromium\*6, if chromium is found in the metals analysis.

Mineral oil is also being considered a potential soil contaminant at all outfalls along the Tijeras Arroyo due to a recent release (June '94) of mineral oil at Outfall 232 and vague historical records. Therefore, soil samples will also be analyzed for TPH.

At Sites 229 through 234, prior to sampling

- 1. tumbleweeds will be cleared from locations to be sampled and placed adjacent to the drainage;
- 2. these locations will be visually scanned for UXO/HE; and
- 3. these locations will be screened for surface radiation anomalies.

Site 229 is due east of the footings of the old guard tower and the south "corner" of the TA II fence. It discharges near the top of the embankment through the center of ER Site 45. Access to this site is along the TA II perimeter road. This site is within the TA II testing exclusion zone. The best days to sample are generally Friday, Saturday, and Sunday, when testing ceases. Bruce Berry (telephone 845-8018) must be contacted to gain permission and access to this site. Because this site discharges from TA II, various radionuclides, possibly including tritium, uranium, and plutonium are of concern. Four surface soil and four subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

Site 230 is west of Building 970 in TAIV. A drain pipe discharges into a bowl-shaped concrete structure adjacent to Building 970A. Flow from this structure is directed to a drain and flume located approximately 120 feet further west. The flume carries the flow to a discharge point slightly above the base of the arroyo embankment. Doug Bloomquist (845-7455) must be contacted to ensure that no laser testing is being performed in the area. Four surface soil and four subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

Site 231 is west of Building 970 in TA IV. A drain pipe discharges to a concrete flume near the top of the embankment. The flume carries the flow to a discharge point near the base of the slope. Doug Bloomquist (845-7455) must be contacted to ensure that no laser testing is being performed in the area.. Four surface soil and four subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

Site 232 consists of two outfalls. One outfall is south of Building 970A, east of the lined lagoon. A drain pipe discharges to a concrete flume near the top of the embankment. The flume carries the flow to at discharge point near the bottom of hillside. On June 1, 1994, about 150 to 350 gallons of mineral oil was spilled into this outfall through the storm water drain by building 986. The day after the spill the site was screened for radiation and UXO/HE. No surface radiation anomalies or UXO/HE were found. Also, four surface soil and four subsurface soil samples were collected. The samples were sent to Quintera Laboratory in Denver for analysis for organics, metals, chromium<sup>+6</sup>, and gamma spec. Other than TPH from the mineral, no contaminants were detected. A Voluntary Corrective Measure was conducted in July and August to remove soil contaminated with mineral oil above 100 mg/kg of TPH.

The second outfall in Site 232 also is south of Building 970A, west of lined lagoon, and approximately 120 feet east of the other Site 232 outfall. Discharge occurs from a concrete structure opening near base of embankment. Access to the site is along the road outside the south side of TA IV. Four surface soil and four subsurface soil samples will be collected at this drainage Figure 1). The analytes are listed in Table 1.

Site 233 is south-southwest of Building 986. Near the top of an escarpment, a small metal drain pipe discharges to an open drain which directs flow within another pipe before discharging near the base of the hillslope. Access to the site is along the road outside the south side of TA IV. Four surface soil and four subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

Site 234 is southeast of Building 981I (Inflatable Building) and a lagoon impoundment (Site 77).

The site discharges into a steep-sided, deeply incised channel cut into the hillside. The drainage channel splits directly uphill of a tree. Access to the site is along the road outside the south side of TA IV. Both channels will be sampled. Six surface soil and six subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

Site 235 is immediately downstream of a large concrete spillway on the northeast side of Pennsylvania and south of the Skeet Range, at the point where the road comes off the north bank of the arroyo and descends into the channel. The flow moves in a confined channel after dropping down the spillway. The site has been cleared for visible surface UXO/HE and screened for surface radiation with no anomalies detected. This channel is considerably larger than the other outfall sites. Six surface soil and six subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

# **Background**

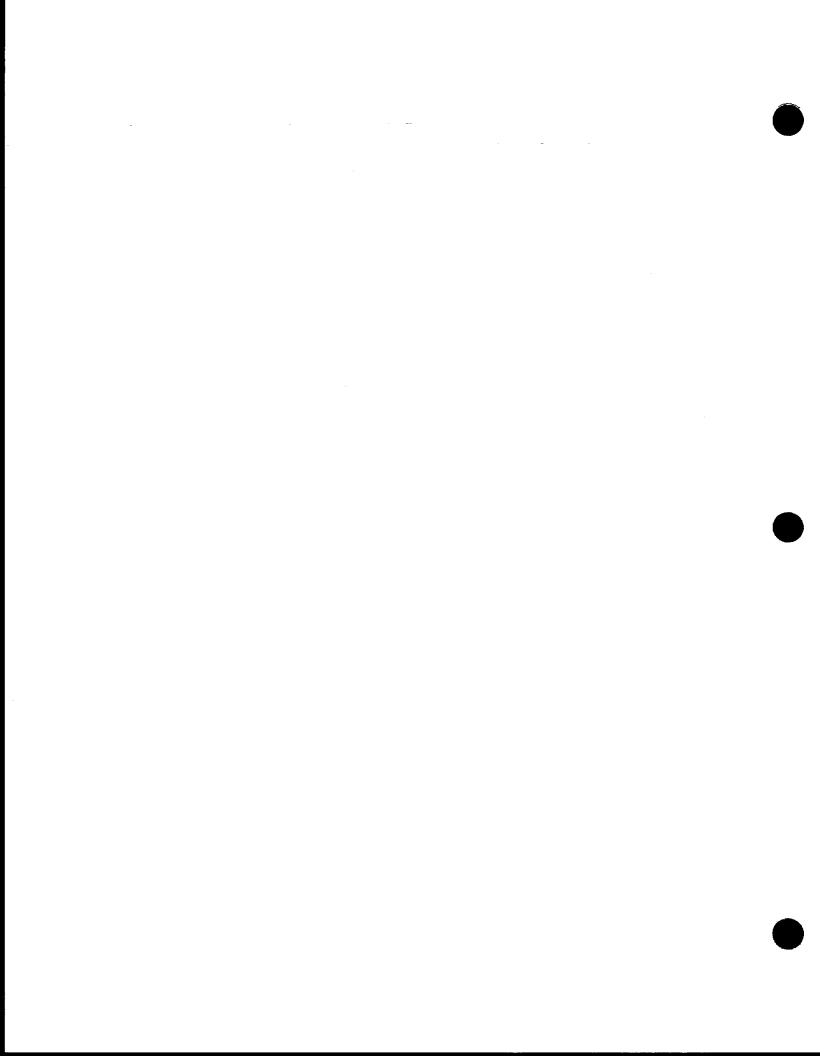
Background soil concentrations for organic contaminants should be negligible. Background concentrations for total metals and radionuclides must be determined for comparison to concentrations found at the sites. Twelve locations have been identified to collect samples for background determination (Figure 1). At each of these sites, one sample will be collected at a depth of 0-6 inches and a second sample collected at 18-36 inches (Table 1).. In addition, the background study report prepared by International Technology Corporation (May 1994) will also be used to evaluate the data.

# **Quality Assurance**

As shown in Table 1, quality assurance samples will include the following:

- Field "duplicates" on more than 10 percent of the samples. These samples will be collected adjacent to the original surface soil sample and in the same hole as the original subsurface soil sample;
- Field soil blanks for more than 10 percent of the VOC analyses. These sample will be obtained from Sample Management Office (SMO) and will contain no VOCs; and
- One rinsate blank. All rinsate will be composited in one container. A sample of the
  rinsate will be analyzed for all constituents. The disposal method for the rinsate will be
  determined by the analytical results on this sample.

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	Number of Samples	4	4	4	4	4	4	4	4	4	9	4	2			58	116	ls a
		and other acids, ng chemicals and hemicals	nd residues	PCBs	e materials, nitrate, solvents, carbon nydroxide, barium, titanium, cyanide	thromium, sodium id, chromosulfuric bleum products	thromium, sodium id, chromosulfuric oleum products	hromium, sodium id, chromosulfuric oleum products	hromium, sodium id, chromosulfuric eleum products	hromlum, sodium d, chromosulfuric leum products	hromium, sodium d, chromosulfuric leum products	nromium, sodium d, chromosulfuric leum products						is detected in meta
	Potential Contaminants	refit chloride, chromic acid and other acids, ammonia, photo processing chemicals and other unknown chemicals	Rocket propellant and residues	Solvents and PCBs	High explosives, radioactive materials, nitrate, toluene, methanol, other solvents, carbon tetrachloride, ammonium hydroxide, barium, cadmium, silver, chromium, titanium, cyanide	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products		S. Y.	a N	Totals	Totals - Surface Plus Subsurface	* Analyze for Cr* only if Cr is detected in metals analysis
	Site Name	Old Acid Waste Line Outfall (Tijeras Arroyo)	Old Centrifuge Site (TA-2)	Oil Surface Impoundment	Bldg. 904 outfall (TA-2)	Storm Drain System Outfall	Background	Duplicates Ciold Seil Brook	Rinsate		Totals - St	*						
Γ	Site		3 2	-+-	227	229	230	=	232	<u></u>	4	ا ور	m .	+	-	$\vdash$	-	j
Ц,	i <u></u> ii	' '		1		73	. 7	231	23	233	234	235	g S	8 8	5 8			



# Appendix B Analytical Results

### ACRONYMS FOR ANALYTICAL DATA

Organic/metals data for soil = mg/kg Radionuclides data for soil = pCi/g

ND = Not detected

NS = Not significant

MDA = Maximum Detectable Activity

J = Detected at a concentration below the laboratory reporting limit

B = Detected in the associated blank sample

Site 46 Soil Results

						_													
	muibo	s	097	250	220	26.4	2 6	270	072	270									
	ilver	s	2 3		2 2			2 2		0.6	462-muines	) O	0.5			α C	5 6	<u>;</u>	
	muinələ	s	2 2				2 2	2 2		3	362\255.muines\	0.02	0.02			0 03	200		
	muisssto	d	000	2500	1500	050	1000	1100	3	301	8&S-muinerL	0.51	0.54		1	0.68	┿	┺	
	јскеј	N	7.0	2   -	7.3	2 0	4 4	4.9	2 2	;	862-muinotul <sup>c</sup>	0.0	00.0			00.0	+-	1	
	Nercury	V	2 2	S	S	S	S	2	S	QN.	04S\eES-muinotul	<0.008	<0.006			<0.004	<0.008		
	ysudsuese	V 6	2 6	210	140	170	160	170	190	200	muitirT	0.04	0.04	0.04	0.04	0.02	0.04	0.03	0.17
	muisangeN	3700	3500	3900	3600	2500	2300	2600	3000	2000	412-beəd					0.93			
IIS	bsə.	1	13			1	7.7	7.8	27	,	212-bead					6.0		-	
SOII RESUITS	ron	12000	13000	17000	12000	8800	11000	8900	11000		04-muisseto9	16.4				22.3			
ñ o t	Copper	) [2	=	12	10	œ	9	11	13		8SS-muibsA	S	QN	-	Q.		9		2
olle o	Cobalt	5.2		5.2	4.3		2.9	3.4	4	.]	822-muibeA	$\vdash$	0.8	-	0.7		0.7		0.7
	muimordO	8.3	15	13	7.7	3.8	5.2	4.4	9.5		მSS-muibeA	SN	NS	NS	2.7	SN	2.1	SN	2.1
	muiɔleɔ	45000	42000	32000	74000	26000	23000	27000	35000		NO3/NO2	1000	1230	1300	1200	1400	150	410	Q.
		ب. 1	4	3.5	2.2	1.5	1.9	1.6	2.7		ТКИ	3.10	88	360	470	120	130	160	190
	Beryllium	0.4	4.0	0.5	0.4	0.2	ΩN	0.2	0.2		əbineyƏ	g	0.5	QN O	QN	9	S	S	0.2
	muineB	180	180	200	220	140	160	140	150		Cr+6*	2	9	9	QN	일	일	2	S.
L	Arsenic	6.7	6.3	7.5	6.7	4.3	-	QN	0.8		Zinc	22	65	2	59	54	21	64	61
	ynomitnA	11	13	17	16	9	8	7	æ		muibeneV	55	97	34	27	138	2	2	77
	munimulA	7000	7900	11000	9500	3700	4200	4400	4500			1	7	T	7	7	$\dashv$	$\dashv$	S
	Sample Identifier	46-01-A	46-01-B	46-02-A	46-02-B	46-03-A	46-03-B	46-04-A	46-04-B		saititnabl alqmaS	46-01-A	46-01-B	46-02-A	46-02-B	46-03-A	46-03-B	46-04-A	46-04-B

Concentrations in mg/kg
Activities in pCi/g
Sample Identifier XX-XX-A - surface soil samples
Sample Identifier XX-XX-B - subsurface soil samples

Quality Assurance Results for Organic Constituents

Нал		T	Τ		T	T	2	2 6	ō		T	$\neg$	$\top$	T	T	T	T	T		Т		2
ofal-Xylenes	1																			1		0000
Styrene									1										- -	+	1	카
Рутеле	0.040 J						1010	0000					1				+	-	+	+		-
Phenanthrene	1=		2				0 17	7 2 2	2						-		†			-		-
Methylene Chloride								1									-		0 003	2		-
Fluoranthene	0.066 J	0.038 J					0.23.1	0.00									-			<u>'</u>	-	-
Di-n-octyl phthalate							-					1	0.16.1					-		-	-	-
Сһтуѕепе							0.11 J	0.12 J										<del> -</del>		1		
Benzo(b)fluoranthene							0.16 J															
Benzo(a)pyrene							0.050 J	0.092														
Benzo(a)anthracene							0.071 J	0.006														
Acetone				0.006													0.019	0.015			0.015	0.010
anonstnaq-S-lydtaM-A			0.001														0.002 J					
S-Hexanone																	0.003 J (	   	<u> </u>			
S-Butanone			0.007 J	0.006 J	0.004 J	0.005 J			0.006 J	0.006	0.006 J	0.006 J	0.003 JB		0.006 JB	0.004 JB	0.010 B	0.009 JB	0.004 JB	8C 200.0	0.007 JB	0.005 JB
Sample Type	original	duplicate	original (	duplicate	original   (	duplicate (	trip blank	trip blank 0	trip blank 0	trip blank 0	trip blank   0	rinsate 0										
Sample Identifier		_	227-01-B		227-04-B	_	229-01-A	-	229-02-B		229-03-B	229-03-B	230-04-B	230-04-B	$\Box$	235-02-B c	-	Site 229   tr		_	$\vdash$	Site 235

		Qualit	y Ass	surand	ce Res	sults fo	or Inord	ranic a	and Rad	noloib	ical Co	netit	uant	_			
Sample Identifier	Sample Type	Aluminum	Antimony	Arsenic	Barium	Beryllium	Cadmium	Chromium	Cobalt	Copper	ron	Lead	Manganese	Mercury	Nickel	Vanadium	Zinc
227-02-A	<del></del>	5800	<del></del>	5.9	180	ND	2.1	6.6	4.1	7.8	13000	7.5	160	ND	5.4	27	51
227-02-A			11	1.4	150	0.25	2.5	6.4	4.1	13	14000	9.1	170	ND	5.9	28	51
227-03-B 227-03-B		5100		0.92	140	ND	2.1	5.9	4.5	11	13000	7.5	200	ND	5.4	25	48
229-04-A				5.6	140	0.25	2.9	7.4	4.6	10	16000	8.9	230	ND	5.9	33	50
229-04-A		8100		5.7	150	0.32	2.3	8.0	4.2	7.9	13000	12	210	ND	6.3	24	55
230-04-B		7700 1500		1.5	140	0.30	2.2	8.0	4.2	7.7	12000	11	190	ND	6.2	24	52
230-04-B				1.6	130	ND	0.61	2.3	ND	18	3500	4.2	110	ND	3.0	9.1	82
235-01-A		3600		1.7	140	ND	0.68	3.1	2.5	15	4500	4.1	120	ND	3.4	9.7	71
235-01-A			6.2 5.3	5.1	150	ND	2.7	6.0	8.4	6.6	20000	7.6	210	ND	4.5	36	66
50-01-B	original	3100		1.3	160	ND	1.6	4.2	5.7	6.5	12000	9.4	180	ND	4.4	22	66
50-01-B	duplicate		6.5	2.1	110	0.25	1.3	4.1	3.9	6.2	7600	6.6	130	ND	4.5	17	18
50-01-B	original	3900	7.5	2.0	110	0.26	1.3	4.3	4.0	5.7	8800	5.9	150	ND	4.2	18	21
50-02-A	duplicate	5800	12	4.2	220	0.38	1.6	5.2	4.3	12	6700	25	210	D	7.1	11.	69
Bkg-05-A		7000	14	6.4	280	0.55	2.2	8.3	6.1	17	9000	35	290	0.04	9.4	18	61
Bkg-05-A	original duplicate	6400	13	5.7	210	0.53	1.8	6.1	6.6	14	10000	16	330	ND	8.9	22	37
Site 235	rinsate	5900	12	7.6	190	0.50	1.7	6.0	6.3	14	10000	16	320	ND	8.7	24	36
C//C 200	moate	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND

Sample Identifier	Sample Type	TKN	NO <sub>3</sub> /NO <sub>2</sub>	Potassium 40	Lead 212	Lead 214	Plutonium 239/240	Uranium 238	Uranium 235/236	Uranium 234
227-02-A	original	400	2.7							
227-02-A		320	9.3							<del></del>
227-03-A	3						0.004	0.4	0.15	0.61
227-03-A								0.67	0.023	0.67
227-03-B	original							0.72	0.11	0.72
227-03-B	original	220	ND							V
227-03-B	duplicate			27.8	0.71	0.7		-		
227-03-B	duplicate	190	1.4						-	
229-01-A	original						0.007	0.45	0.17	0.67
229-01-A	duplicate							0.73	0.034	0.6
229-03-B	original							0.45	0.058	0.45
229-03-B	duplicate							0.99	0.06	1

### Notes on Quality Assurance Data

Explosive residues were not detected in Site 50 duplicate sample

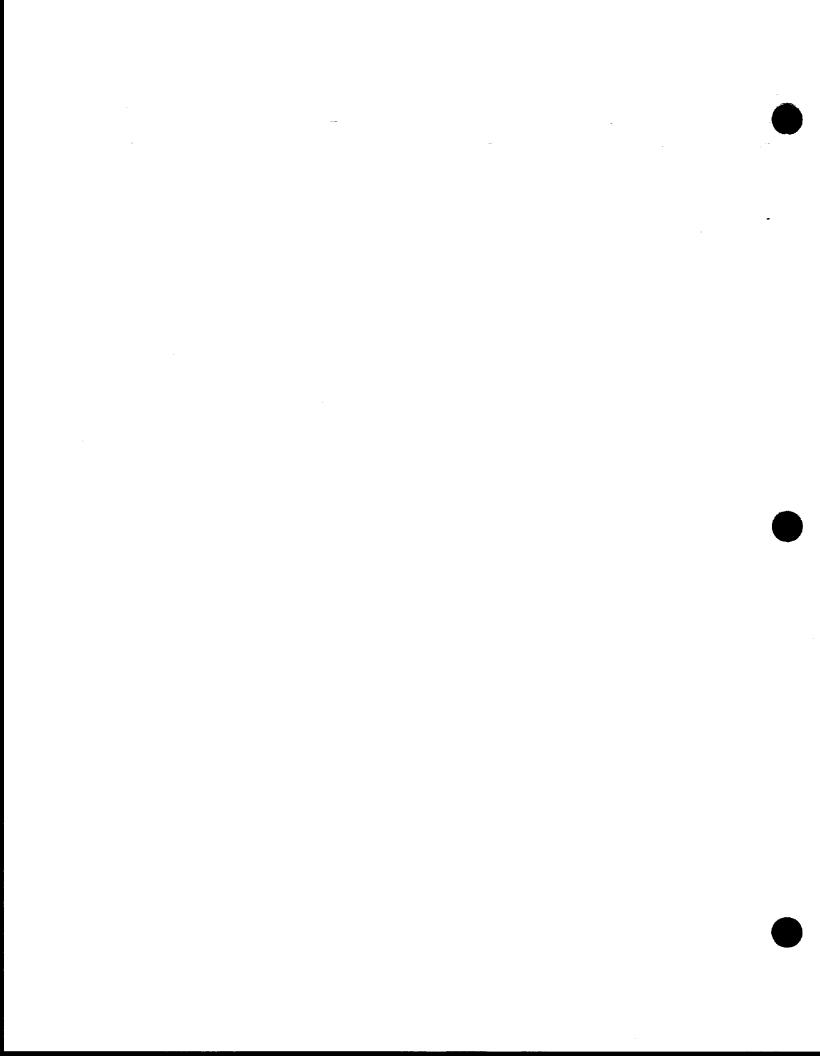
Hexavalent chromium was not detected in five duplicates and one decon rinsate

Cyanide was not detected in two duplicates and one decon rinsate

PCBs were not detected in one Site 77 duplicate sample

Tritium and Plutonium-238 were not detected in four duplicate samples

Selenium, silver, and thallium were not detected in any quality assurance samples



### Appendix C Background Calculations for Metals and Radionuclides

### Appendix C. Background Calculations for Metals and Radionuclides

To evaluate metals data, 24 background samples were collected for metals analyses.<sup>4</sup> Distribution analyses was performed first by constructing histograms. The histograms indicated a parametric distribution. Outliers were screened in a two-step process as described in the base wide background report (IT 1994). The first step is to perform an "a priori" screening for very high values relative to the rest of the data set. This is qualitatively performed by visually examining a column of sorted values. Maximum values that are a factor of 3 or 4 times higher than their nearest neighbor are removed from the data set during this step. None of the anomalous values were deleted by the "a priori" process.

The second step, from EPA, 1989, determines whether an observation that appears extreme fits the data distribution. A statistical parameter,  $T_n$  is calculated:

$$T_n = (X_n - X_n)/S$$

where:

 $X_n$  = questionable observation;

 $X_a$  = sample arithmetic mean; and

S = sample standard deviation

 $T_n$  is compared to a table of one-sided critical values for the appropriate significance level (upper 5 percent) and sample size from a table provided in EPA 1989. Extreme concentrations for barium, calcium, chromium, copper and nickel were identified as outliers and were excluded from the data set. These anomalous values may have resulted from laboratory or sampling error.

Probability plots were then replotted to determine whether the data fit normal or lognormal populations. These plots are shown in Appendix D. The UTL<sup>5</sup> was calculated for data sets that fit a normal or lognormal distribution. Data sets are provided in Appendix D. As recommended by EPA, a tolerance coefficient value of 95 percent was used (EPA 1989). Most metals background data fit lognormal distributions. Iron and zinc data fit normal distributions. UTLs were not calculated for mercury, selenium, and silver because mercury and selenium were not detected and silver was detected only once in the 24 background samples. The beryllium background data did not fit a normal or lognormal distribution. The maximum value in a data set is commonly taken as the UTL in a non-parametric setting (Guttman, 1970). The maximum background beryllium concentration was 0.53 mg/kg.

Base-wide background UTLs for radionuclides were established by International Technology (IT) Corporation to compare and evaluate radionuclide data (IT, 1994). A table is provided in Appendix

<sup>&</sup>lt;sup>2</sup>These data are referred to as local background data. The data collected throughout Kirtland Air Force Base (KAFB), with most of the data collected within SNL/NM technical areas, are called base-wide background data (IT 1994).

 $<sup>^3</sup>$ UTL = x + K•S, where:

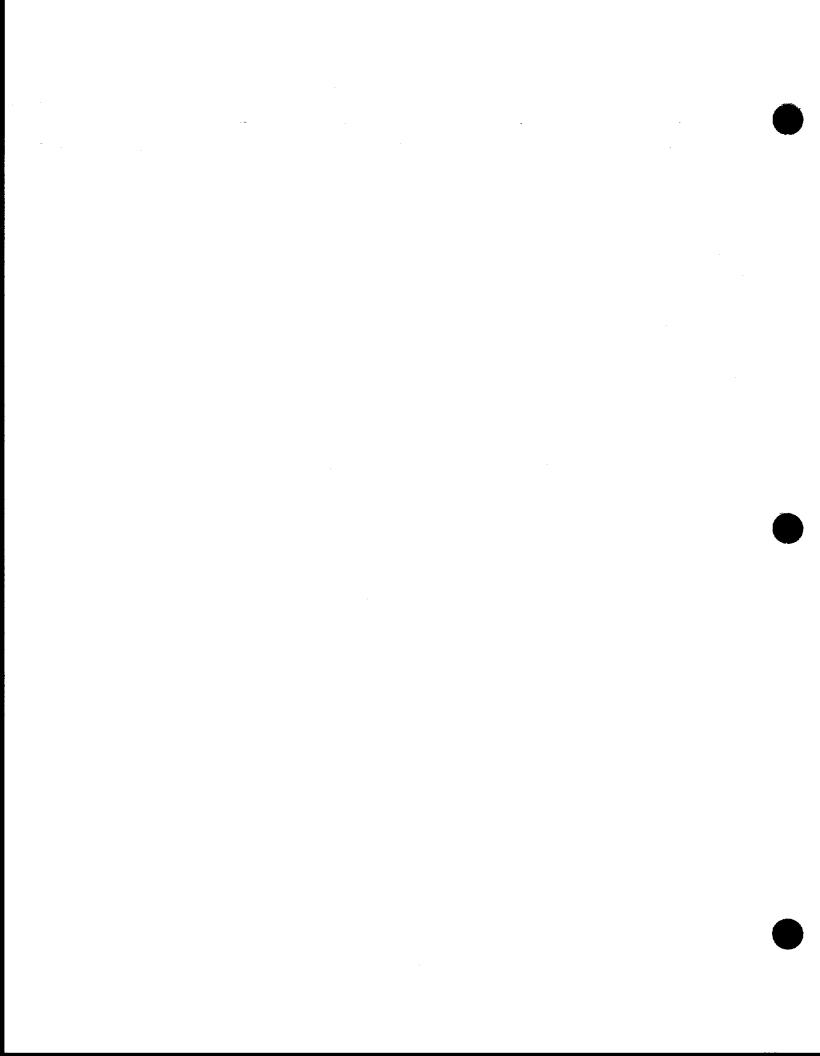
UTL = Upper tolerance limit;

x = Sample arithmetic mean (for normal distribution), sample geometric mean (for lognormal distribution);

S = Sample standard deviation; and

K = One-sided normal tolerance factor (95 percent for these evaluations).

D with radionuclide background data and the corresponding UTLs. The maximum activity from the six local background samples for isotopic plutonium and isotopic uranium was used as an additional method to evaluate the data. Also, in-house gamma spectroscopy was performed on all 24 background samples and indicated low levels of radioactivity but no significant contamination.

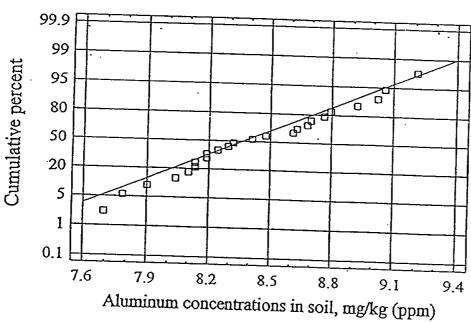


## Appendix D Probability Plots, Local Background UTL Calculations, and BaseWide Background UTLs for Radionuclides

Summary Statistics for log(Aluminum)

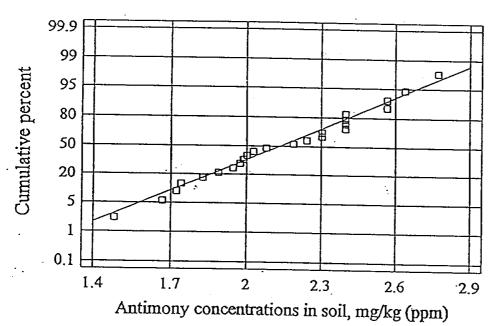
Count = 24 Average = 8.42942Median = 0.36529Mode = Geometric mean = 8.41976 Variance = 0.170246 Standard deviation = 0.412609 Standard error = 0.0842235 Minimum = 7.69621 Maximum = 9.21034 Range = 1.51413 Lower quartile = 8.13153 Upper quartile = 8.73178 Interquartile range = 0.600253 Skewness = 0.132255 Stnd. skewness = 0.26451Kurtosis = -0.792361Stnd. kurtosis = -0.792361Coeff. of variation = 4.89487 Sum = 202.306

### Lognormal Probability Plot for Aluminum



Summary Statistics for log(Antimony) .verage = 2.14609 edian = 2.13275 ode = 2.3979 cometric mean = 2.12004 ariance = 0.113831 tandard deviation = 0.337389 tandard error = 0.0688692 inimum = 1.4816 aximum = 2.77259 ange = 1.29098ower quartile = 1.91649 oper quartile = 2.3979 iterquartile range = 0.481405 cewness = -0.040772ind. skewness = -0.0815441ictosis = -0.744171ind. kurtosis = -0.744171 peff. of variation = 15.7211 m = 51.5062

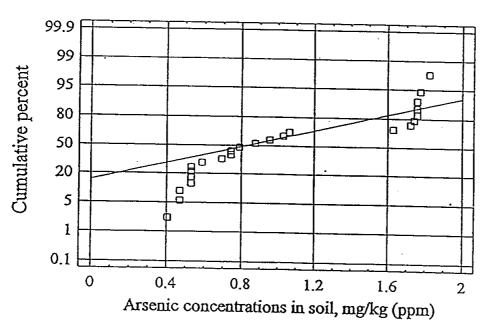
### Lognormal Probability Plot for Antimony



Summary Statistics for log(Arsenic)

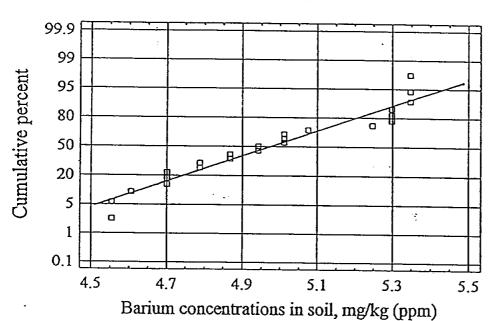
Count = 24 Average = 1.038 Median = 0.031963Mode = Geometric mean = 0.908119 Variance = 0.291153 Standard deviation = 0.539586 Standard error = 0.110143 Minimum = 0.405465 Maximum = 1.82455 Range = 1.41908 Lower quartile = 0.530628 Upper quartile = 1.73162 Interquartile range = 1.20099 Skewness = 0.463036 Stnd. skewness = 0.926071{urtosis = -1.58507  $\exists$ tnd. kurtosis = -1.58507 Doeff. of variation = 51.983 3um = 24.9121

### Lognormal Probability Plot for Arsenic



ry Statistics for log(Bacium) ecage = 4.96948 dian = 4.94164de = 5.34711ometric mean = 4.96236 ciance = 0.0740602 andard deviation = 0.27214 andard error = 0.0567451 mium = 4.55388 <imum = 5.34711 ige = 0.793231 ver quartile = 4.70048 per quartile = 5.29832 erquartile range = 0.597837 wness = 0.0653415d. skewness = 0.127931tosis = -1.30542 d. kurtosis = -1.27794ff. of variation = 5.47622= 114.298

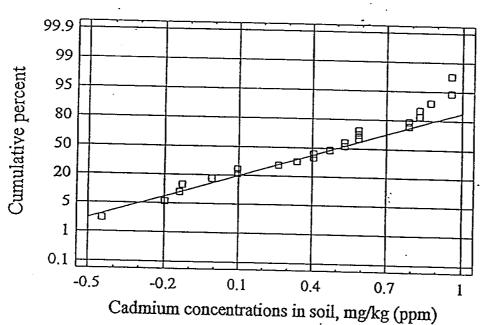
### Lognormal Probability Plot for Barium



Summary Statistics for log(Cadmium)

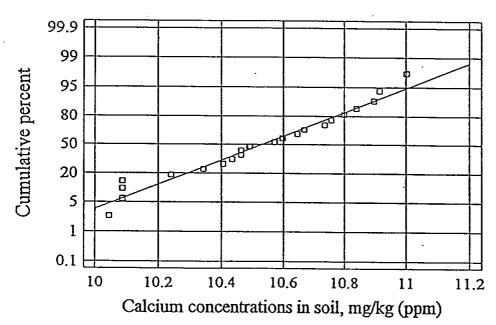
Count = 24 Average = 0.416764 Median = 0.500316 Mode = Geometric mean = Variance = 0.159937 Standard deviation = 0.399922 Standard error = 0.0816337 Minimum = -0.446287Maximum = 0.955511 Range = 1.4018 Lower quartile = 0.0953102 Upper quartile = 0.788457 Interquartile range = 0.693147 Skewness = -0.506707Stnd. skewness = -1.01341Kurtosis = -0.674504 Stnd. kurtosis = -0.674504Coeff. of variation = 95.9587 Sum = 10.0023

### Lognormal Probability Plot for Cadmium



Jummary Statistics for log(Calcium) ge = 10.5579dian = 10.5713 x = 10.0058ometric mean = 10.5532 riance = 0.10513andard deviation = 0.324237 andard error = 0.0676081 nimum = 10.0432 |ximum = 11.2645 inge = 1.22121 wer quartile = 10.3417 per quartile = 10.7996 terquartile range = 0.457833 ewness = 0.109797 nd. skewness = 0.214971rtosis = -0.415646nd. kurtosis = -0.406895 eff. of variation = 3.07103 m = 242.832

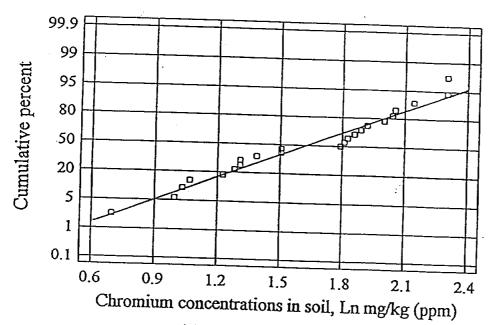
### Lognormal Probability Plot for Calcium



Summary Statistics for log(Chromium)

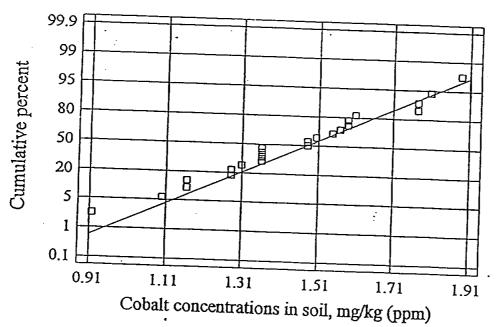
Count = 23Average = 1.61841 Median = 1.79176 Mode = Geometric mean = 1.55042Variance = 0.204195 Standard deviation = 0.451879 Standard error = 0.0942233 Minimum = 0.693147Maximum = 2.30259 Range = 1.60944 Lower quartile = 1.28093 Upper quartile = 2.00148 Interquartile range = 0.720546 Skewness = -0.274151Stnd. skewness = -0.536757Kurtosis = -0.905395 Stnd. kurtosis = -0.886332Coeff. of variation = 27.9211Sum = 37.2235

### Lognormal Probability Plot for Chromium



ary Statistics for log(Cobalt) Average = 1.29969 Median = 1.42129Mode = Geometric mean = /ariance = 0.574775 Standard deviation = 0.758139 Standard error = 0.154754 linimum = -2.07944faximum = 1.88707 tange = 3.96651ower quartile = 1.28093 oper quartile = 1.58924 nterquartile range = 0.308301 kewness = -4.13299tnd. skewness = -8.26598urtosis = 10.9091 tnd. kurtosis = 18.9091 oeff. of variation = 58.3324 um = 31.1925

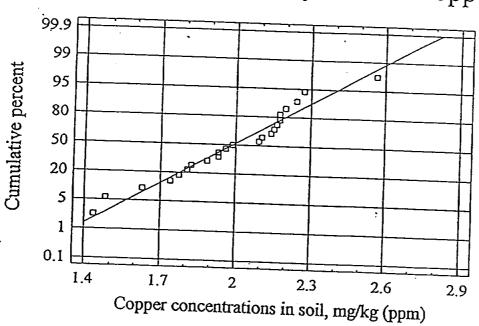
### Lognormal Probability Plot for Cobalt



Summary Statistics for log(Copper)

Count = 23 Average = 1.98556 Median = 1.98787 Mode = Geometric mean = 1.96762 Variance = 0.0713494 Standard deviation = 0.267113 Standard error = 0.0556969 Minimum = 1.43508 Maximum = 2.56495 Range = 1.12986 Lower quartile = 1.80829 Upper quartile = 2.17475 Interquartile range = 0.366463 Skewness = -0.263077 Stnd. skewness = -0.515077Kurtosis = 0.18883 Stnd. kurtosis = 0.184854 Coeff. of variation = 13.4528 Sum = 45.6679

### Lognormal Probability Plot for Copper

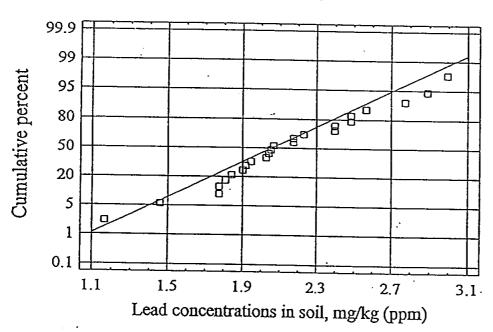


Summary Statistics for log(Lead)

ge = 2.13936 Median = 2.06049 Mode ⊨ Geometric mean = 2.09509 Variance = 0.187882 Standard deviation = 0.433454 Standard error = 0.0884784 Minimum = 1.16315 Maximum = 2.99573 Range = 1.83258Lower quartile = 1.87133 Jpper quartile = 2.4414 Interquartile range = 0.570072 kewness = 0.0350174 itnd. skewness = 0.0700348urtosis = 0.200156 itnd. kurtosis = 0.200156'oeff. of variation = 20.261

um = 51.3446

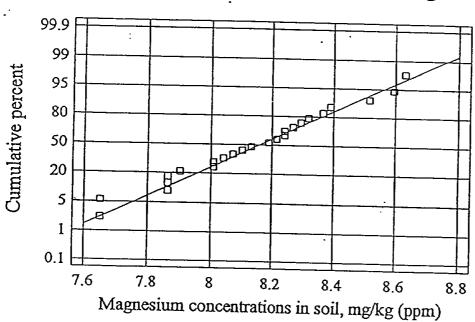
### Lognormal Probability Plot for Lead



Summary Statistics for log(Magnesium)

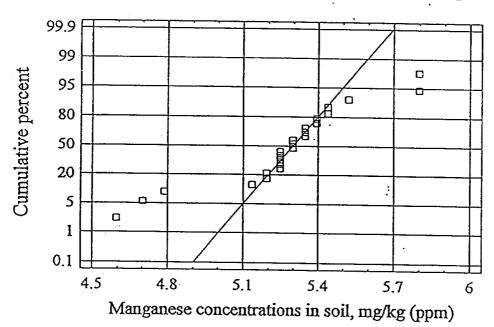
Count = 24Average = 8.14232 Median = 8.16011 Mode = Geometric mean = 8.13815 Variance = 0.0706013 Standard deviation = 0.265709 Standard error = 0.0542376 Minimum = 7.64969 Maximum = 8.63052 Range = 0.980829 Lower quartile = 7.95369 Upper quartile = 8.3064 Interquartile range = 0.352709 Skewness = -0.0600481Stnd. skewness = -0.120096Kurtosis = -0.414246 Stnd. kurtosis = -0.414246Coeff. of variation = 3.26331 Sum = 195.416

### Lognormal Probability Plot for Magnesium



Summary Statistics for log(Manganese) rage = 5.2733 Median = 5.29832 Mode = Geometric mean = 5.2661 Variance = 0.0771874 Standard deviation = 0.277826 Standard error = 0.056711 Minimum = 4.59512 Maximum = 5.79909 Range = 1.20397Cower quartile = 5.21999 Jpper quartile = 5.39363 interquartile range = 0.173637 3 kewness = -0.6603873tnd. skewness = -1.32077(urtosis = 1.62566 Stnd. kurtosis = 1.62566 Soeff. of variation = 5.26854 um = 126.559

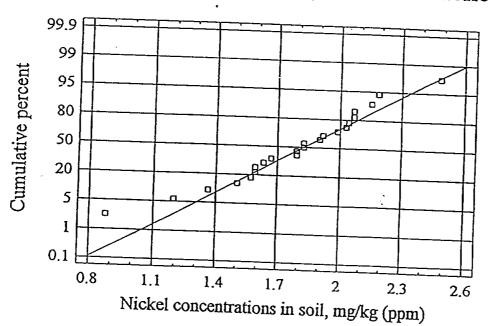
### Lognormal Probability Plot for Manganese



Summary Statistics for log(Nickel)

Count = 23 Average = 1.78451Median = 1.82455Mode = Geometric mean = 1.74596 Variance = 0.1246 Standard deviation = 0.352987 Standard error = 0.0736029 Minimum = 0.875469Maximum = 2.48491 Range = 1.60944 Lower quartile = 1.58924 Upper quartile = 2.04122 Interquartile range = 0.451985 Skewness = -0.609856Stnd. skewness = -1.19403Kurtosis = 0.992502 Stnd. kurtosis = 0.971605 Coeff. of variation = 19.7806 Sum = 41.0438

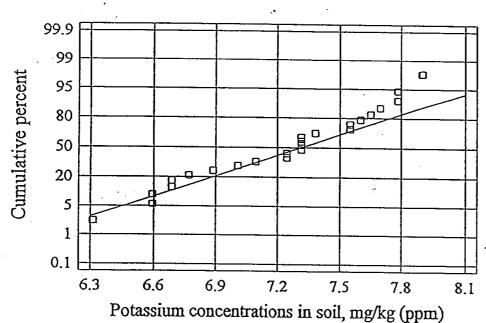
### Lognormal Probability Plot for Nickel



immacy Statistics for log(Potassium)

= 7.21862 :dian = 7.31322 ide = 7.31322ometric mean = 7.20542 .ciance = 0.195599andard deviation = 0.442265 andard error = 0.0902771 nimum = 6.30992ximum = 7.90101nge = 1.59109 wer quartile = 6.82802 per quartile = 7.57526 terquartile range = 0.747233 ewness = -0.373735 nd. skewness = -0.74747ctosis = -0.83864nd. kurtosis = -0.83864eff. of variation = 6.12673 n = 173.247

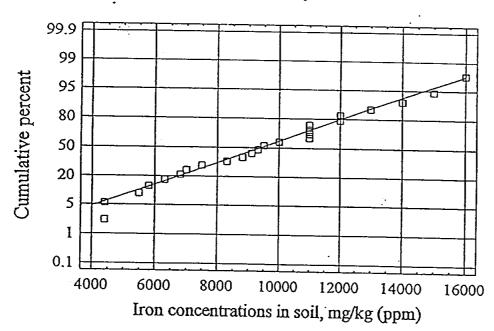
### Lognormal Probability Plot for Potassium



Summary Statistics for Iron

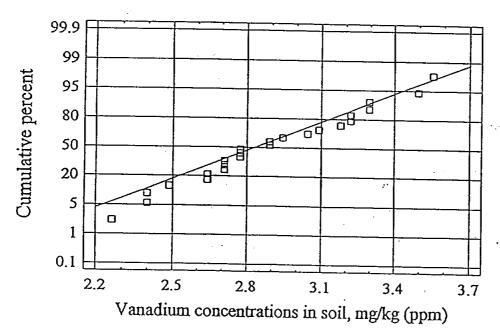
Count = 24 Average = 9529.17 Median = 9400.0 Mode = 11000.0Geometric mean = 8977.5 Variance = 1.0363E7 Standard deviation = 3219.17 Standard error = 657.109 Minimum = 4400.0 Maximum = 16000.0 Range = 11600.0 Lower quartile = 6900.0 Upper quartile = 11500.0 Interquartile range = 4600.0 Skewness = 0.20025Stnd. skewness = 0.400499Kurtosis = -0.620589Stnd. kurtosis = -0.620589Coeff. of variation = 33.7822 Sum = 228700.0

### Normal Probability Plot for Iron



ummary Statistics for log(Vanadium) /erage = 2.89094 edian = 2.83148ode = ometric mean = 2.87064 rlance = 0.122444 andard deviation = 0.34992 andard error = 0.0714271 nimum = 2.26176 ximum = 3.55535 nge = 1.29358wer quartile = 2.67355 per quartile = 3.19846 terquartile range = 0.524911 ewness = 0.158415 nd. skewness = 0.316831rtosis = -0.688491 nd. kurtosis = -0.688491eff. of variation = 12.104 m = 69.3826

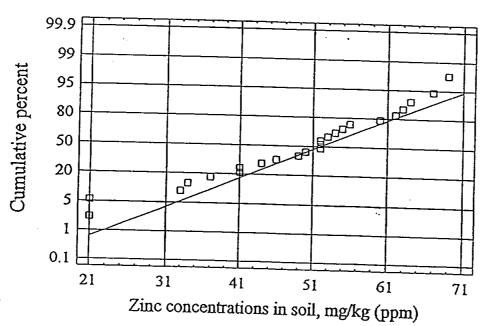
### Lognormal Probability Plot for Vanadium



Summary Statistics for Zinc

Average = 49.0 Median = 52.0Mode = 52.0Geometric mean = 46.9434 Variance = 171.478 Standard deviation = 13.095 Standard error = 2.673 Minimum = 21.0 Maximum = 69.0 Range = 48.0Lower quartile = 41.0 Upper quartile = 58.0 Interquartile range = 17.0 Skewness = -0.633044Stnd. skewness = -1.26609Kurtosis = -0.0224531 Stnd. kurtosis = -0.0224531Coeff. of variation = 26.7244Sum = 1176.0

### Normal Probability Plot for Zinc



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	ysuðsuese	۸ <u>۶</u>	2 6	3 6	210	210	250	190	210	330	330	190	200	170	180	110	180	230	120	190	200	061	220		220
	muisəngsN	2100	3100	2100	2700	3700	4400	3000	2600	5600	3900	3800	3400	2600	3000	2600	4100	5400	3800	3200	3300	-		4300	
	bea	ع اد	1	ო	ω	0	တ	8	စ	16	-	8	7	12	7	4	9	8	9	12	11	8		၈	13
	uou	5800	8800	4400	6300	11000	11000	9300	8300	10000	11000	9100	6800	7000	7500	4400	9500	11000	5500	13000	14000	12000	16000	12000	15000
	Copper	) ဖ		4	6	တ	တ	13	7	14	6	8	7	9	9	4	7	7	2	8	9	6	6	6	6
	tlado	) m	4	က	က	ည	ည	4	4	7	9	4	4	4	4	ΩN	4	ည	3	2	4	ည	9	2	9
	muimondC	) m	2	7	က	9	7	6	4	9	9	19	4	4	4	3	5	6	3	8	7	8	0,	7	5
	Calcium	23000	24000	35000	31000	36000	28000	24000	24000	78000	33000	46000	51000	34000.	39000	54000	00009	49000	82000	42000	35000	43000	40000	55000	47000
	muimbe	0.9	1.5	9.0	1	1,8	1.8	2.3	1.4	1.8	1.7	1.5	1.1	1.1	1.3	9.0	1.6	1.7	6.9	2.3	2.6	2.2	2.4	2.2	2.6
_	Beryllium	9	0.3	DN	ND	0.4	0.4	0.3	ND	0.5	0.5	0.3	0.3	0.3	0.3	원	2	4	0.3	0.3	0.3	0.4	0.5		0.4
_	Rarium	110	130	110	130	110	92	120	120	210	40	150	150	95	2	<u>8</u>	190	210	210	140	150	200	200		290
L	Arsenic	7	2	7	7	2	7	7	7	9	9	9	7	7	က	0	က	٥	70	7	9	7	7	7	9
L	ynomitnA	9	8	4		6	의		9	13	9	6	8	9	7	ω l	1	= -	7	=	=	3	16	=	4
	munimulA	2700	4100	2400	3400	4800	0009	4000	3300	6400	5500	4500	3800	3100	3600	2200	3000	2900	3400	7500	0099	8300	10000	2600	8600
	Sample Identifier	Bkg-01-A	Bkg-01-B	Bkg-02-A	Bkg-02-B	Bkg-03-A	BKg-03-B	BKg-04-A	вкд-04-в	Bkg-05-A	Bkg-05-B	BKg-06-A	Bkg-06-B	Bkg-07-A	Bkg-07-B	BKg-08-A	BKG-08-B	BKg-09-A	a-80-Bya	BKg-10-A	Bkg-10-B	Bkg-11-A	Bkg-11-B	Bkg-12-A	Bkg-12-B

Concentrations in mg/kg
Activities in pCi/g
Sample Identifier XX-XX-A - surface soil samples
Sample Identifier XX-XX-B - subsurface soil samples

Local Background Soil Results

	462-muine							-	. 0	?					T							C C	90	0.8	0.9
	1ranium-235/236							0.28	000	2010												0.03	0.03	0.17	0.33
	862-muiner(							80	0										1					<del>                                     </del>	0.6
	86S muinotul	<u> </u>						< 0.011	<0.00													<0.017	<0.018	< 0.017	0.038
21	₽S\882 muinotul	1						<0.009	<0.008													<0.007	<0.012	<0.030	0.035
and provided the police	muitinT							<0.010	<0.022													<0.023	<0.024	<0.084	<0.023
	oni <u>S</u>	2	63	41	53	99	62	55	52	37	34	52	54	21	21	33	67	4	44	52	49	09	64	46	69
	muibensV	=	16	9.6	=	19	15	18	16	22	18	16	14	15	15	12	21	24	14	27	27	25	35	25	33
<u> </u>	muilledT	S	2	Q	Q.	N N	2	2	2	2	Q.	QΝ	S	9	2	2	9	9	2	2	S	身	2	QN	N N
	muiboS	2	S	ND	Q.	2	480	2	욷	9	2	ΔN	420	9	380	<u>Q</u>	430	280	640	Q.	S	280	_	QN	620
	19vli2	Q N	Q	ND	QN	g	g	+	QN	QN	Q.	QN	QN	2	S	g	Q.	QN	ND	9	S	S	QN	ΔN	QN
	Gelenium	g	ND	ΔN	ND	QΝ	ND	QN	QN	QN	ND	N N	QN N	9	Q.	QΝ	ND	QΝ	QN	QΝ	9	Q.	ND	QN	QN
	muisssto9	1500	2000	730	1600	1500	1200	1900	1400	2700	1400	1500	800	870	800	730	980	1100	550	2400	2200	2100	2400	1500	1900
	Nickel	4	9	7	2	7	6	12	ည	6	8	13	9	5	5	က	ည	ω	2	9	7	7	8	9	<u></u>
	Sample Identifier	Bkg-01-A	Bkg-01-B	Bkg-02-A	Bkg-02-B	Bkg-03-A	Bkg-03-B	Bkg-04-A	Bkg-04-B	Bkg-05-A	Bkg-05-B	Bkg-06-A	Bkg-06-B	Bkg-07-A	Bkg-07-B	Bkg-08-A	Bkg-08-B	Bkg-09-A	Bkg-09-B	Bkg-10-A	Bkg-10-B	Bkg-11-A	Bkg-11-B	Bkg-12-A	Bkg-12-B

Concentrations in mg/kg
Activities in pCi/g
Sample Identifier XX-XX-A - surface soil şamples
Sample Identifier XX-XX-B - subsurface soil samples

Normal Parameters for Tijeras Arroyo Local Metal Background Data Aluminum Antimony Manganese Cadmium Monadium 17 Statistical Cobalt Nickel Lead **Parameter** ron median 4300 8.5 2 140 2 6 4.2 7.3 9400 geometric mean 7.9 200 4579.9 52 8.6 144 2 5 3.7 7.3 8977.5 maximum 8.5 195 6 10000 18 47 16 6 210 3 10 6.6 13 16000 20 minimum 330 12 35 2200 4.4 69 2 95 1 2 0.1 4.2 arithmetic average 4970.8 4400 3.2 99 2.4 9.6 9 21 3 149 2 5.5 4.2 7.5 9529.2 standard deviation 2095.4 9.3 202 6.3 19 49 3 2 40.5 1 2.3 1.3 2 normal tolerance 3219.2 4.2 53.6 2.1 6.9 2.309 2.3 2 13 2.33 2 2.3 2.3 2.3 2.309 2.3 2.31 UTL 4927.4 2.3 2.3 16 2.3 244

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Log	normal F	aram	eter	s for T	iiera	c Arr	ovo i	1							1
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	uminum	Š			Ę	mium					sse		_		I
Statistical	i j	ţi	eni:	n nic	dmic	ļ <u>Ē</u>	జ	per	·		gane	<b> </b>	Enil	1	ļ
Parameter	1 7	5	Arse	Bari	Cad	Chro	go	d	5	ad	ang	ckel	anadi	ပ	
rithmetic average	8.4294	2.2	1	4.97	) 0	1.6	1.3	ာ	<u> </u>		Σ	ž	S	Zinc	ĺ
tandard deviation	0.4126	0.3	1	0.27	0	0.5			9.1025		5.27	1.8	2.9	3.8	
normal tolerance	2.309	2.3	2	2.33	2	2.3	0.8	0.3	0.3631	0.4	0.28	0.4	0.3	0.3	
UTL	9:3821	2.9	2	5.6		2.7	2.3	2.3	2.309	2.3	2.31	2.3	2.3	2.3	
e <sup>UTL</sup>	11874	19	10	271	<del>-    </del>		3.1	2.6	9.941	3.1	5.91	2.6	3.7	4.6	
			.01	2/1	4	14	21	14	20764	23	370	14	40	98	

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insufficient data for mercury, selenium, silver, and thallium to calculate statistics All concentrations in mg/kg-

# Summary of Background Concentrations for Radionuclides in Soil

95" Upper Tolerance Limit 95" Percenile	_		/''	9.0	-	0.92	- «detection limit	(<0.0686)	Adetection limit	(<0.041B)	6.8	565	-		- 1	0.568	1	50.5		0.300	+	- 8			0.	0.168	
	(pCI/g) (pC	1.0	90	0.0		. 0.2495	<detection limit<="" p=""></detection>	(<0.0000)	(<0.0418)		2.835	0.5 1.0795	0.56	+	10.4	0.655	0.590	0.630	'	0.2883	+	1	0.71 2.89	1	6.0	0.1235	
Geometric Mean	(bc/g)	1,1055	0.648	2000	1 ,	0.200		1	<u> </u>	0.0000	4,40030	0.49689	0.549	15.889		0.6747	0.713	0.695		0.2528	0.7971	1/2/0	08770	t	0.897	0,1198	
	e	13	321		1 8	50 t		247	; ;	48	,	233	240	718	7	\$	54	2	0	45	136	35	;	3	4	8	
Ranga	(Rand)	0.414-2.7	0.27-1.4		1 60 0	Adetection limit	(<0.0686)	-detection limit	(<0.0418)	0.3-12.0		0.1-1.4	0.29-1.13	0.192-31.0	0 49 0 07	/Ein-orio	0.5-2.09	0.45-1.05	ŧ	0.0321.85	0.23-1.20	0.324~3.0		1	0.6-8.0	81.0=-c0.0	- 0000
Distribution	N	Nonparametric	Nonparametric		Nonparametric	Unknown		Unknown	• *	Nonparametric	Language	2011001100	Lognormal	Normal	Nonoarametric	Lograna	Cognormal	Nonparametric	Unknown	Nonparametric	Lognormal	Lognormal	Unknown	olstom essential	Monogrammenic	Nonparament	
Number of Rejected Samples	307		19	28	1	,		74		292	06		2	*	0	314		5   6	0	6	0	330	0	0	75	1,2	-
Number of Detects	11	305	351	561	ı	ı		=	,	40	233	241		/20	24	83	24		,	45	136	52	0	4	21	206	-
Originat Number of Samples	324	340		802	:	1		321	338	000	323	249	100	77,	24	368	22		13	5   5	927	365	0	4	95	223	_
Analyte	Bismuth-212	Bismuth-214	Casium, 127	(Surface)	(Subsurface)	(Social ace)	Coballieo	00.1800	Lead-210°		Lead-212"	Lead-214°	Potassium.40		Madium-224	Radium-228	Radium-228	Radon	Strontium.an	Thorium.232	1	1 norium-234	Trilium	Uranium-234	Uranium-235	Uranium-238	

'Sample size. 'Sample size. \*These constituents are not listed as COC in Table 2-2 for this media. \*Constituents of concern are of unknown distribution type because data are either below the limit of detection, unusable, or nonexistent.

(IT, 1994)

The second secon